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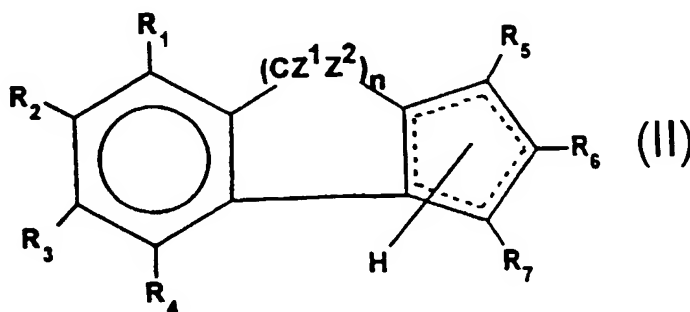
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- (74) Agents: DE GREGORI, Antonella et al.; Ing. Barzanó & Zanardo Milano S.p.A., Via Borgonuovo, 10, I-20121 Milan (IT).
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- (71) Applicant (*for all designated States except US*): ENICHEM S.P.A. [IT/IT]; Piazza Boldrini, 1, I-20097 San Donato Milanese (IT).
- (72) Inventors; and
- (75) Inventors/Applicants (*for US only*): BIAGINI, Paolo [IT/IT]; Via Gramsci, 73, I-28069 Trecate (IT). VIGLIAROLO, Diego [IT/IT]; Via A. Ratti, 37, I-20017 Rho (IT). BORSOTTI, Giampietro [IT/IT]; Strada Pastore, 14, I-28100 Novara (IT). SANTI, Roberto [IT/IT]; Viale Piazza d'Armi, 24/F, I-28100 Novara (IT).
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(54) Title: SUBSTITUTED POLYCYCLIC CYCLOPENTADIENES AND METHOD FOR THEIR PREPARATION



(57) Abstract: New polycyclic cyclopentadiene compounds having the formula (II) wherein the various substituents and symbols R_1 , R_2 , R_3 , R_4 , R_5 , R_6 , R_7 , Z^1 and Z^2 and "n" have the meaning specified in the description. These compounds can form metallocene complexes with transition metals, which have shown unusual properties in the (co)polymerization of ethylene and alpha-olefins in general.

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SUBSTITUTED POLYCYCLIC CYCLOPENTADIENES AND METHOD FOR
THEIR PREPARATION.

The present invention relates to compounds of the
10 group of cyclopentadienes substituted to form polycyclic
structures, as well as a process for their preparation.

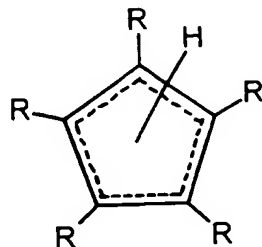
More specifically, the present invention relates to
new polycyclic cyclopentadienyl compounds which can be
used, in the form of anions, as penta-hapto (η^5 -) ligands
15 coordinated to transition metals to form metallocenes.

The property of cyclopentadienyl compounds of easily
forming an anion of an aromatic nature by the loss of a
cationic fragment, usually an H^+ anion, is known. The ca-
pacity of these anions of coordinating themselves to metal
20 cations by means of a multi-electronic complex bond to form
complexes, called metallocenes, having an unusual relative
stability with respect to the behaviour generally expected
for organometallic compounds containing hydrocarbon anions,
is also well known. These complexes may contain a single
25 cyclopentadienyl ligand, or 2 and, in certain cases, even

three ligands of this type. Cyclopentadienyl complexes of transition metals of Groups 3 to 6, and particularly of Group 4 of the periodic table, combined with aluminoxanes or particular salts of non-coordinating anions, are known to be capable of polymerizing α -olefins and have been widely used for this purpose in the recent past.

Cyclopentadienyl compounds comprise the di-unsaturated cyclic group with five carbon atoms having the following general formula (I):

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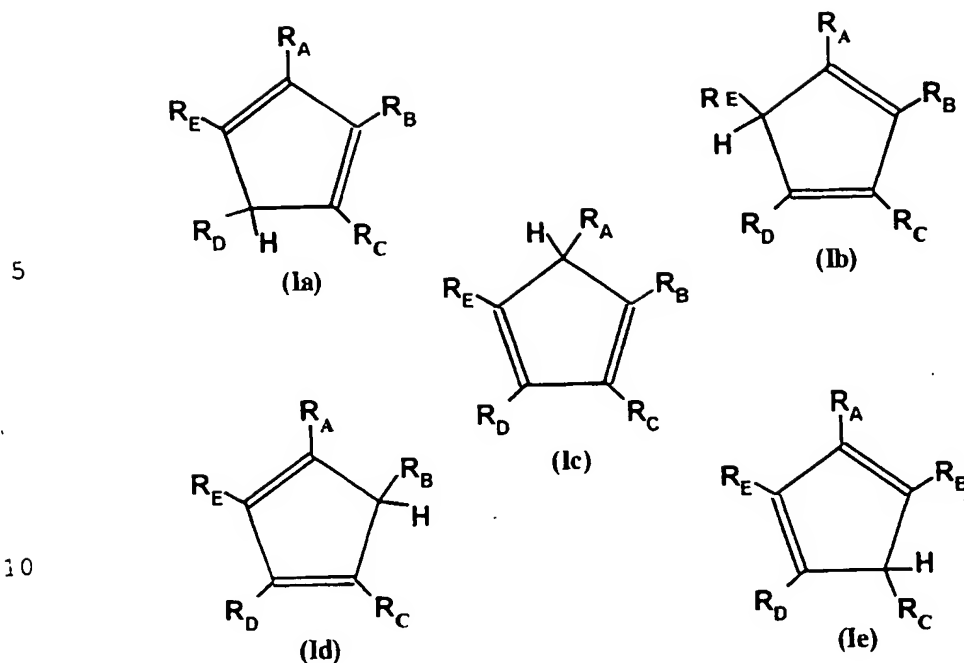


(I)

wherein the R groups represent general substituents, either the same or different.

In accordance with the present invention and in order to simplify the description and claims, the above formula (I) is used herein to synthetically represent one or more of the five possible mesomeric structures (Ia), (Ib), (Ic), (Id) and (Ie) indicated below, which, as is normally customary in organic chemistry, can represent a general polysubstituted cyclopentadienyl compound.

25



These can be converted to one another by the shifting, in the pentagonal cycle, of the two double bonds and the hydrogen atom linked to the sole saturated carbon atom, the various substituents of the cycle usually being hydrogen or linear or branched organic radicals, optionally substituted, which, in the more general version, can also form one or more covalent ligands with each other to obtain cyclic or polycyclic structures condensed with said cyclopentadienyl ring. For the purposes of the present invention, these structures (Ia), (Ib), (Ic), (Id) and (Ie) can be considered as being equivalent, as they are precursors of the same cyclopentadienyl anion. Consequently, unless otherwise specified, the scope of the present invention, with

25

reference to the cyclopentadienyl compounds represented by structural formulae of the type (I), which form one of the objects thereof, comprises the five mesomeric structures defined above.

- 5 Substituted cyclopentadienyl hydrocarbon compounds consisting of saturated and unsaturated rings condensed on the cyclopentadienyl ring are known in the art, and some of them are commercial products, such as, for example, indene, tetrahydroindene, fluorene, azulene, 1,2-benzofluorene.
- 10 Other bicyclic compounds comprising a cyclopentadienyl ring are described in European patent application EP-A 760,355, whereas European patent application EP-A 849,273 describes metallocene complexes with these ligands, suitable for polymerizing olefins.
- 15 The positive effect of saturated cycles condensed with cyclopentadienyl rings in the polymerization catalysis of olefins, using metallocenes of zirconium and titanium, has also been acknowledged, especially with respect to the activity and incorporation of co-monomers in the co-
- 20 polymerization of ethylene with α -olefins, as described for example in European patent application EP-A 849,273.

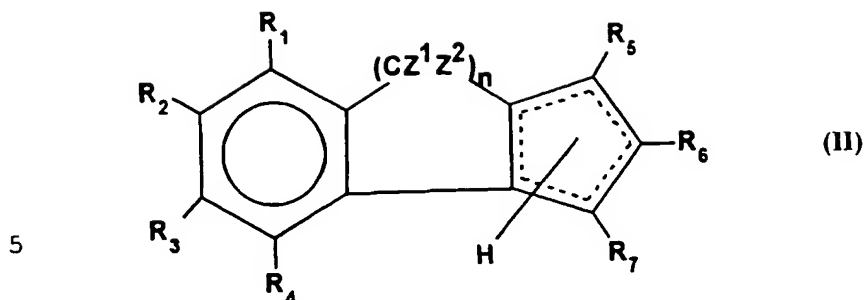
In spite of the encouraging results obtained so far in this field, there is still the request for continuous improvements in polymerization catalysts and in the quality

25 and versatility of poly- α -olefins obtained therewith. The

request for elastomeric polymers of ethylene with a high content of unsaturations suitable for a rapid vulcanization, has stimulated research relating to catalysts capable of increasing the quantity of non-conjugated dienes incorporated in the ethylene copolymer with respect to what is so far possible, without causing a significant decrease in the average molecular weight of the product obtained. However it is desirable to have catalysts capable of producing polyolefins with an increased molecular weight for high yield processes operating at temperatures higher than 100°C. Finally, further improvements are required in the steric control of the stereospecific polymerization of α -olefins to give polymers with a high tacticity by means of metallocene catalysts of the so-called "bridged" type, having structures which are becoming more and more sophisticated and selective.

In the never-ending attempt to respond to the above demands with innovative processes and materials, the Applicant has now found a new group of polycyclic cyclopentadienyl compounds, suitable for use in the preparation of catalysts which represent a definite progress in providing a solution to the above problems.

A first object of the present invention therefore relates to a polycyclic cyclopentadienyl compound having the following formula (II):



wherein: each R_1 , R_2 , R_3 , R_4 , R_5 , R_6 , R_7 , Z^1 and Z^2 is independently hydrogen or an organic substituent having from 1 to 10 carbon atoms, preferably selected from hydrocarbon radicals having from 1 to 8 carbon atoms, more preferably C_1 - C_8 alkyl, linear or branched, and in addition, any one of said R or Z groups, preferably one of the groups selected from Z^1 , Z^2 , R_5 , R_6 , or R_7 , can be a divalent organic group further bound to another organic group having from 5 to 20 carbon atoms and comprising a cyclopentadienyl group, and

"n" has any of the integer values from 1 to 10 extremes included, and preferably ranges from 1 to 3.

As already specified above, in the above formula (II), the cyclopentadienyl group is conventionally represented by a structure which comprises all the mesomeric forms obtainable by shifting the double bonds and the hydrogen atom in the cycle as required by the valency regulations. This

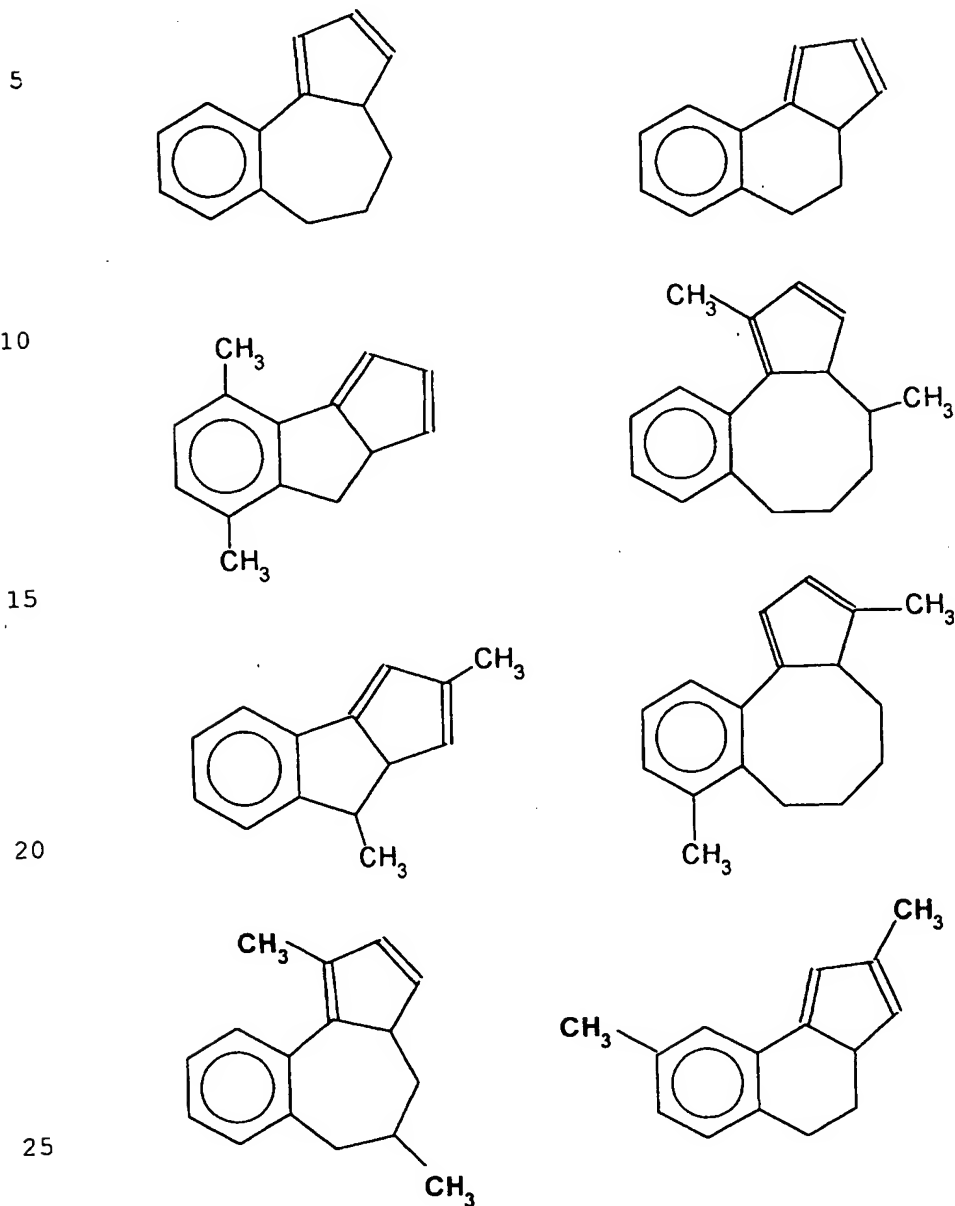
shifting is not only formal, but can be easily obtained in practice by forming a cyclopentadienyl anion of any compound having formula (II), for example by treatment with lithium butyl, and by subsequently acidifying to obtain a
5 mixture of all possible five isomers, said mixture being also included in the scope of the present invention.

In a preferred embodiment, the present invention relates to compounds having formula (II) wherein all the R and Z groups are hydrogen. In another preferred embodiment,
10 Z^1 and Z^2 are hydrogen and only two of the groups R_1 , R_2 , R_3 , R_4 are linear or branched alkyl with from 1 to 4 carbon atoms.

According to other particular forms of the present invention, one of the R_5 , R_6 or R_7 groups is methyl or ethyl,
15 more preferably methyl. In another preferred embodiment, R_1 and R_3 are methyl, and "n" is 2 or 3.

According to a particular embodiment of the present invention, one of the R_5 , R_6 or R_7 groups, preferably R_5 or R_7 , is a divalent hydrocarbon or silane group having from 2
20 to 6 carbon atoms, further linked to a second cyclopentadienyl group having from 5 to 20 carbon atoms, preferably selected from cyclopentadienyl, indenyl, fluorenyl and homologous groups thereof, in particular linked to said R_5 , R_6 or R_7 group by means of a carbon atom of the cyclopenta-
25 dienyl ring.

Non-limiting examples of compounds having formula (II) are those represented by the following formulae, which for the purposes of simplicity, are not numbered:

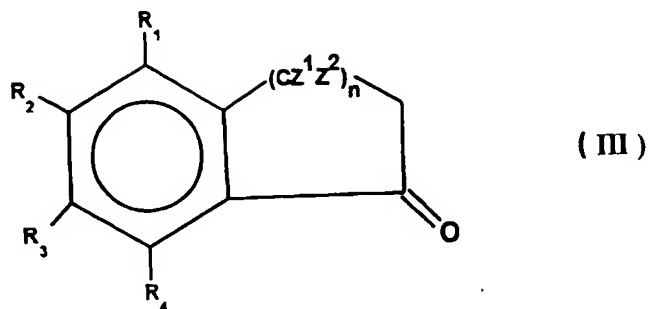


The polycyclic cyclopentadienyl compounds according to the present invention are relatively stable compounds, they can be exposed to air, even if it is not convenient to keep them for long periods due to the possibility of the formation of the corresponding dimeric products, according to the well known Diels-Alder reaction. For this reason, when there is the possible necessity of keeping these products for some time, it is preferable to transform them into one of the corresponding salts of alkaline or earth-alkaline metals, which, although being sensitive to oxygen and humidity present in the air, do not undergo any chemical alterations if kept in an inert atmosphere.

These polycyclic compounds, although new, can be prepared by adapting known techniques of organic chemistry for the purpose, by analogy with synthetic methods of compounds and organic structures already known or similar, such as, for example, the synthesis method described in the European patent application cited above EP-A 760,355. The Applicant however has developed an original synthetic method, by modifying or combining known techniques, which allow these compounds to be obtained with a good yield in a limited number of steps.

A second object of the present invention therefore relates to a method for the preparation of the polycyclic cyclopentadienyl compounds having formula (II) starting

from a benzoketone having the following general formula
(III):

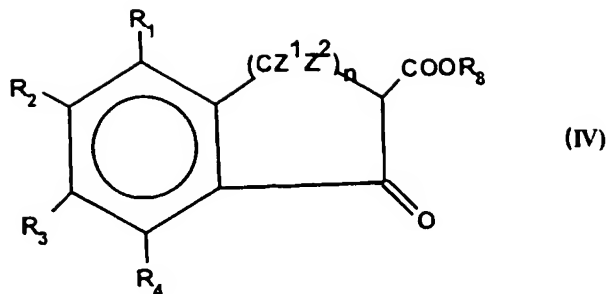


wherein the various symbols have the same meaning defined above, both in the general and in the preferred forms, with reference to the cyclopentadienyl compounds having formula (II),

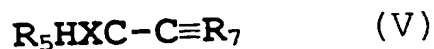
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comprising the following steps in succession:

- i) forming the anion of said benzoketone having formula (III) by the extraction of a hydrogen ion in alpha position with respect to the carbonyl, by means of reaction with a hydride or an alkyl of an alkaline metal,
- 15 ii) reacting the anion formed in step (i) with an alkyl carbonate having the formula $OC(OR_3)$ wherein R_3 is linear or branched alkyl having from 1 to 6 carbon atoms, to obtain the benzoketoester having the following
- 20 formula (IV):

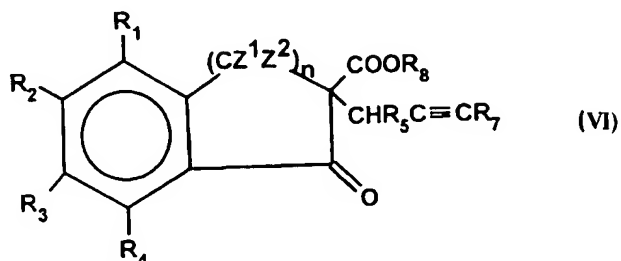


- iii) forming the anion of said benzoketoester having formula (IV) by reaction with a sufficiently strong organic base, for example a hydride, an alkoxide or an alkyl of an alkaline metal, and reacting said anion with a propargyl halide having the following formula (V)
- 10



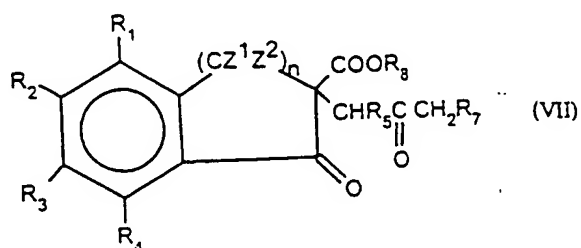
- 15 wherein R_5 and R_7 have the same meaning as the corresponding symbols of formula (II) and X is halogen excluding fluorine, preferably chlorine or bromine, to obtain at the end the following compound having formula (VI)

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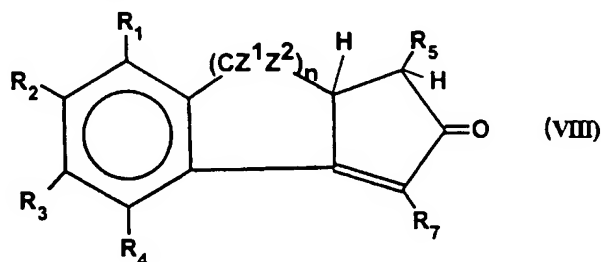


- 25 iv) hydrating said compound having formula (VI) with one

of the known methods for transforming the acetylene group into a ketone group, for example by contact and reaction in a mixture with mercury oxide, etherate boron trifluoride, trichloroacetic acid in methanol as solvent, and subsequent hydrolysis, in order to obtain the following compound having formula (VII):

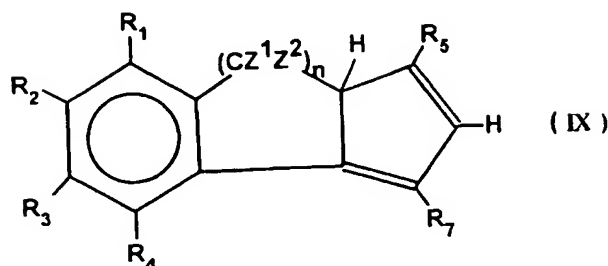


- v) cyclizing said compound having formula (VII) by self-condensation in a strongly basic environment by KOH or NaOH, at temperatures preferably ranging from 20 to 100°C, in order to obtain the following unsaturated cyclic ketone having formula (VIII):



- vi) reducing said unsaturated cyclic ketone having formula (VIII) using one of the known methods, for example by reaction with lithium aluminum hydride, to obtain the desired polycyclic cyclopentadienyl compound having

formula (IX), corresponding to the compound having
formula (II) with R_7 equal to hydrogen:



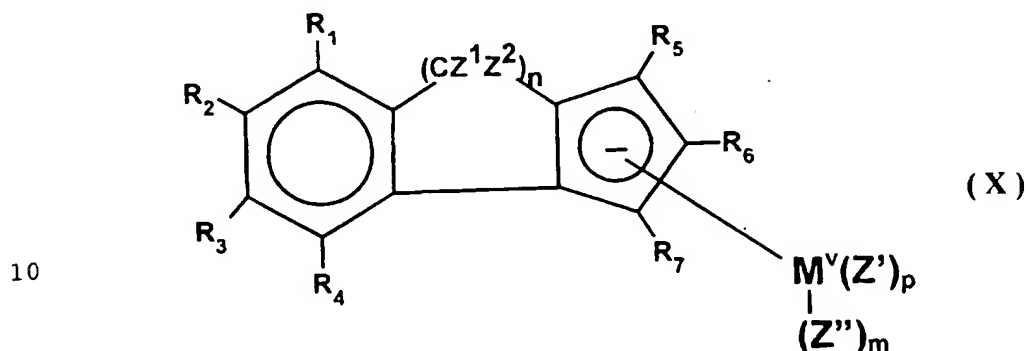
or by means of reaction with a suitable reactive metal
alkyl comprising the R_8 group, for example lithium al-
10 kyl, a Grignard reagent, a magnesium dialkyl, an alu-
minum trialkyl etc., to obtain the desired compound
having general formula (II) object of the present in-
vention.

The above polycyclic cyclopentadienyl compounds can be
15 used for the formation of organometallic complexes of tran-
sition metals, particularly of group 4 of the periodic ta-
ble of elements, which can be advantageously used as cata-
lysts in various synthetic processes, such as for example,
hydrogenation, oxidation and polymerization of olefins.

20 A further object of the present invention therefore
relates to saline (ionic) compounds comprising at least one
polycyclic cyclopentadienyl anion formally deriving from
any of the previous compounds having formula (II) by the
extraction of an H^+ acid anion from the cyclopentadienyl
25 ring, and also metal complexes comprising at least one of

said cyclopentadienyl anions coordinated to a transition metal by means of covalent bonds of the π type. In particular, an object of the present invention relates to an organometallic compound having the following general formula

5 (X):



wherein:

each R_1 , R_2 , R_3 , R_4 , R_5 , R_6 , R_7 , Z^1 and Z^2 group, and also "n", has the same meaning, both general and preferred, as the corresponding symbol in formula (II) above;

M^v represents any metal of the periodic table of elements having an oxidation state (or valence) "v" greater than zero;

each Z' is independently a group of an anionic nature bound to the metal M^v as anion in an ionic couple or with a covalent bond of the " σ " type;

Z'' represents an organic radical having from 5 to 30 carbon atoms, comprising a cyclopentadienyl anion bound to the metal M^v ;

"m" has the value 1 or 0 depending on whether Z" is present or absent in the compound having formula (X);

"p" expresses the number of Z' groups necessary for neutralizing the formal oxidation charge "v" of the metal M^v and has such values that $p = (v - m - 1)$.

The above formula (X) includes all compounds comprising a metal M^v in an oxidation state greater than zero and at least one anion deriving from a ligand having the previous formula (II) according to what is specified above.

10 These compounds can be of a saline ionic nature, especially with non-transition metals, or, when low energy orbitals of the metal are available to form bonds of the "π" type, these compounds are preferably in the form of co-ordinated complexes of a semi-covalent or covalent nature.

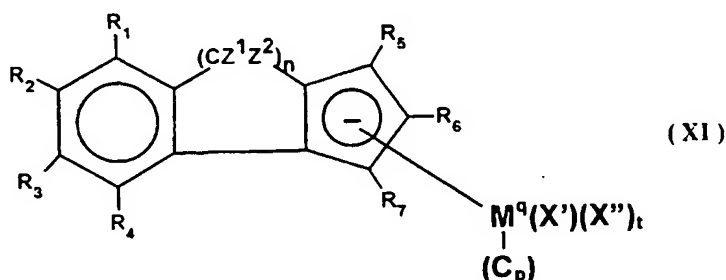
15 Of particular interest for their potential use as catalysts or components of catalysts for hydrogenation or polymerization processes of α-olefins, are compounds having formula (X) wherein M^v is selected from metals of groups 3 to 10 of the periodic table, including metals of the group

20 of lanthanides.

M^v is preferably selected from metals of groups 4 to 6, and is more preferably Ti, Zr or Hf in oxidation state +4. In this case, particularly satisfactory results have been obtained when the Z" group in formula (X) is a second

25 cyclopentadienyl anion, optionally, but not necessarily,

selected from compounds having formula (II) above. In particular, a preferred object of the present invention relates to a bis-cyclopentadienyl complex of a transition metal M^3 of group 4 of the periodic table, having the following formula (XI):



wherein:

each R_1 , R_2 , R_3 , R_4 , R_5 , R_6 , R_7 , Z^1 and Z^2 group, and also "n", has the same meaning, both general and preferred, as the corresponding symbol in formulae (II) and (X) above;

M^3 represents a metal selected from titanium, zirconium and hafnium having an oxidation state (or valence) "q" equal to 3 or 4, preferably 4;

"t" has the value of 1 if the valence "q" of M^3 is 4, and the value of 0 if the valence of "q" of M^3 is 3;

X' and X'' each independently represent a group of a monovalent anionic nature σ -bound to the metal M^3 , preferably selected from hydride, halide, a C_1 - C_{20} alkyl or alkylaryl group, a C_2 - C_{20} alkylsilyl group, a

C_2-C_{10} cycloalkyl group, a C_6-C_{10} aryl or arylalkyl group, a C_1-C_{10} alkoxyl or thioalkoxyl group, a C_2-C_{10} carboxylate or carbamate group, a C_2-C_{10} dialkylamide group and a C_1-C_{10} alkylsilylamide group.

5 C_F represents any organic group containing a η^5 -cyclopentadienyl or η^5 -heterocyclopentadienyl ring, of an anionic nature, preferably having at least from 5 to 30 carbon atoms, coordinated to the metal M^3 .

According to the present invention, in particular, the
10 groups X' and X'' having formula (XI) each independently represent a group of an anionic nature σ -bound to the metal M^3 . Typical examples of X' and X'' are hydride, halide, preferably chloride or bromide, a linear or branched alkyl group such as methyl, ethyl, butyl, isopropyl, isoamyl, oc-
15 tyl, decyl, benzyl, an alkylsilyl group such as, for example, trimethylsilyl, triethylsilyl or tributylsilyl, a cycloalkyl group such as cyclopentyl, cyclohexyl, 4-methylcyclohexyl, an aryl group such as phenyl or toluyl, an alkoxyl or thioalkoxyl group such as methoxyl, ethoxyl,
20 iso- or sec-butoxyl, ethylsulfide, a carboxylate group such as acetate, trifluoroacetate, propionate, butyrate, pivalate, stearate, benzoate, or again a dialkylamide group such as diethylamide, dibutylamide, or alkylsilyl-amide, such as bis(trimethylsilyl)amide or ethyltrimethylsilylamide.
25 The two X' and X'' groups can also be chemically bound to

each other and form a cycle having from 4 to 7 atoms different from hydrogen, also comprising the metal M^I . Typical examples of this aspect are divalent anionic groups such as the trimethylene or tetramethylene group, or the ethylenedioxy group. X' and X'' groups which are particularly preferred for their accessibility and the easy preparation of the complexes which comprise them are chloride, C_1-C_3 alkoxide, methyl, ethyl, C_3-C_7 carboxylate.

Each C_5 group in formula (XI) contains a η^5 -cyclopentadienyl ring coordinated to the metal M^3 , which formally derives from a cyclopentadiene molecule, substituted or non-substituted, by extraction of an H^+ ion. The molecular bond of the π type and also the electronic and coordinative configuration of metallocene complexes of titanium, zirconium or hafnium, generally comprising two η^5 -cyclopentadienyl groups, have been amply described in literature and are known to experts in the field.

The C_5 group having formula (XI) according to the present invention is selected, in particular, from known cyclopentadienyl, indenyl or fluorenyl groups, and their homologous products, in which one or more carbon atoms of the molecular skeleton (included or not included in the cyclopentadienyl ring), are substituted with a radical selected from the group consisting of halogen, preferably chlorine or bromine, a linear or branched alkyl group having from 1

to 10 carbon atoms, optionally halogenated, such as methyl, trifluoromethyl, ethyl, butyl, isopropyl, isoamyl, octyl, decyl, benzyl, an alkylsilyl group such as, for example, trimethylsilyl, triethylsilyl or tributylsilyl, a cycloalkyl group such as cyclopentyl, cyclohexyl, 4-methylcyclohexyl, an aryl group having from 6 to 10 carbon atoms, optionally halogenated, such as phenyl, pentafluorophenyl or toluyl, an alkoxyl or thioalkoxyl group such as methoxyl, ethoxyl, iso- or sec-butoxyl, ethylsulfide. Said Cp group may also comprise several condensed aromatic rings, as in the case, for example, of 4,5-benzoindenyl, and in a particularly preferred embodiment, is selected from anions of polycyclic cyclopentadienyl compounds having formula (II) described above.

For particular polymerization processes, especially when the steric control of the polymerization (iso- or syndiotactic orientation of α -olefins) is important, or the control of the insertion of co-monomers in the copolymerization of ethylene, as is known to experts in the field, the above groups of an anionic nature, more specifically Z^- in the complexes having formula (X) with M' a transition metal, and C_p^- in complexes having formula (XI), can be covalently bound to the polycyclic cyclopentadienyl group deriving from compounds having formula (II), by means of a divalent hydrocarbon or silane group having from 1 to 15,

preferably from 2 to 6, carbon atoms, to form a so-called "bridged" structure, according to the traditional terminology used in the field of metallocene compounds.

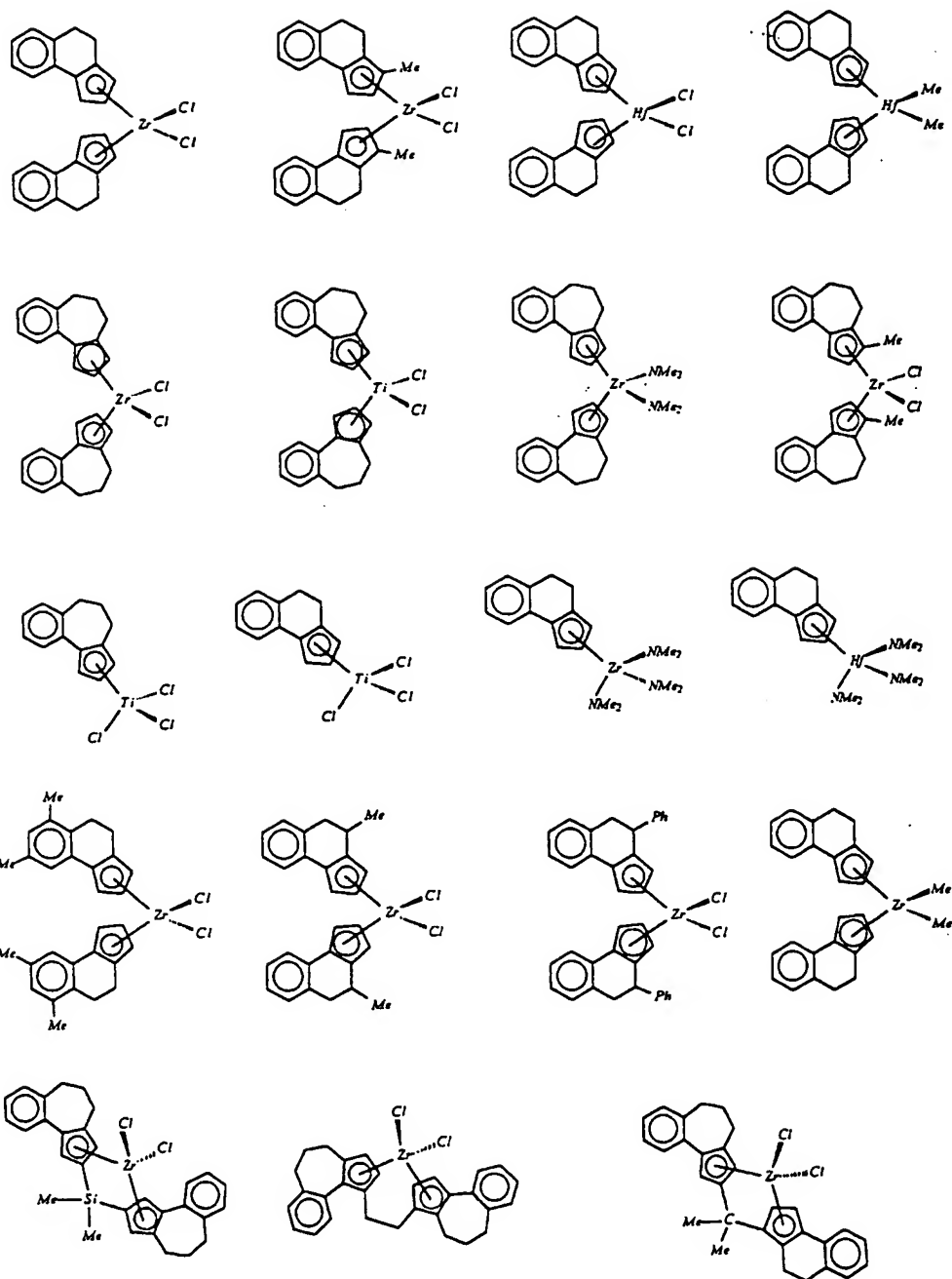
Non-limiting examples of compounds having general formula (X) and (XI) are those represented by the structures indicated hereunder, which, for the sake of simplicity, are not numbered:

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As mentioned above, the above complexes having formula (X) with certain transition metals, and more specifically complexes represented by formula (XI), can be advantageously used in the formation of catalysts for the polymerization and co-polymerization of olefins, combined with a suitable activator.

A further object of the present invention therefore relates to, and is claimed herein as such, a catalyst for the (co)polymerization of ethylene and other α -olefins, i.e. for the homo-polymerization of ethylene and other α -olefins, the co-polymerization of ethylene with one or more other co-polymerizable monomers, such as for example, α -olefins, conjugated or non-conjugated diolefins, styrene and its derivatives, etc., the co-polymerization of α -olefins with each other or with other monomers copolymerizable therewith. This catalyst can be obtained from the combination (i.e. contact and reaction) of the above polycyclic metallocene complex having formula (X) or (XI) with an activator (or co-catalyst) selected from those known in the art of metallocene polymerization catalysis, particularly an organic compound of a metal M' selected from boron, aluminum, gallium and tin, or a combination of these compounds.

In particular, said catalyst in accordance with the present invention comprises the following two components in

contact with each other, or the product of their reaction:

- (i) at least one metallocene complex having the previous formula (X), wherein the metal M'' is a metal of groups 4 to 6, preferably 4, of the periodic table;
- 5 (ii) a co-catalyst consisting of at least one organometallic compound of an element M' different from carbon and selected from elements of groups 2, 12, 13 or 14 of the periodic table as defined above.

According to the present invention, said element M' is
10 preferably selected from boron, aluminum, zinc, magnesium, gallium and tin, more particularly boron and aluminum.

In a preferred embodiment of the present invention, the component (ii) is an organo-oxygenated derivative of aluminum, gallium or tin. This can be defined as an organic
15 compound of M' , in which the latter is bound to at least one oxygen atom and to at least one organic group consisting of an alkyl group having from 1 to 6 carbon atoms, preferably methyl.

According to this aspect of the invention, component
20 (ii) is more preferably an aluminoxane. As is known, aluminoxanes are compounds containing Al-O-Al bonds, with a varying O/Al ratio, which can be obtained, under controlled conditions, by the reaction of an aluminum alkyl or aluminum alkyl halide, with water or other compounds containing
25 pre-established quantities of water available, such as, for

example, in the case of the reaction of aluminum trimethyl with aluminum sulfate hexahydrate, copper sulfate pentahydrate or iron sulfate pentahydrate. Aluminoxanes preferably used for the formation of the polymerization catalyst of the present invention are oligo- poly-meric, cyclic and/or linear compounds, characterized by the presence of repetitive units having the following formula (XII):



10 wherein R_8 is a C_1 - C_8 alkyl group, preferably methyl.

Each dialuminoxane molecule preferably contains from 4 to 70 repetitive units which may also not all be equal to each other, but contain different R_8 groups.

Said aluminoxanes, and particularly methylaluminoxane, 15 are compounds which can be obtained by known methods in organometallic chemistry, for example, by the addition of aluminum trimethyl to a suspension in hexane of aluminum sulfate hydrate.

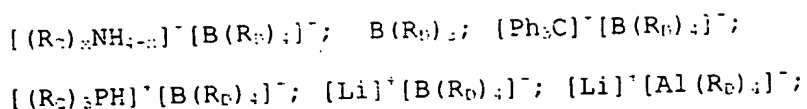
When used for the formation of a polymerization catalyst according to the present invention, the aluminoxanes 20 (ii) are put in contact with said component (i) comprising the complex having formula (X) in such proportions that the atomic ratio between Al and the transition metal M is within the range of 10 to 10,000 and preferably from 100 to 25 5,000. The sequence with which the complex (i) and the alu-

minoxane (ii) are put in contact with each other, is not particularly critical.

In addition to the above aluminoxanes, the definition of component (ii) according to the present invention also comprises galloxanes (in which, in the previous formulae, gallium is present instead of aluminum) and stannoxanes, whose use as cocatalysts for the polymerization of olefins in the presence of metallocene complexes is known, for example, from patents US 5,128,295 and US 5,258,475.

10 According to another preferred aspect of the present invention, said catalyst can be obtained by putting component (i) consisting of at least one complex having formula (X), in contact with component (ii) consisting of at least one compound or a mixture of organometallic compounds of M' capable of reacting with the complex having formula (X),
15 extracting from this a σ -bound group Z' or Z'' to form, on the one hand at least one neutral compound, and on the other hand an ionic compound consisting of a metallocene cation containing the metal M^V and an organic non-coordinating anion containing the metal M', whose negative
20 charge is delocalized on a multicentric structure.

Components (ii) suitable as ionizing systems of the above type are preferably selected from the voluminous organic compounds of boron and aluminum, such as for example,
25 those represented by the following general formulae:



wherein the deponent "x" is an integer ranging from 0 to 3,
 each R_2 group independently represents an alkyl or aryl
 5 radical having from 1 to 10 carbon atoms and each R_2 group
 independently represents an aryl radical partially or,
 preferably, totally fluorinated, having from 6 to 20 carbon
 atoms.

Said compounds are generally used in such quantities
 10 that the ratio between the atom M' of component (ii) and
 the atom M'' in the metallocene complex is within the range
 of 0.1 to 15, preferably from 0.5 to 10, more preferably
 from 1 to 6.

Component (ii) can consist of a single compound, nor-
 15 mally an ionic compound, or a combination of this compound
 with MAO, or, preferably, with an aluminum trialkyl having
 from 1 to 8 carbon atoms in each alkyl residue, such as for
 example $AlMe_3$, $AlEt_3$, $Al(i-Bu)_3$.

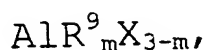
In general, the formation of the ionic metallocene
 20 catalyst, in accordance with the present invention, is
 preferably carried out in an inert liquid medium, more
 preferably hydrocarbon. The selection of components (i) and
 (ii), which are preferably combined with each other, as
 well as the particular method used, can vary depending on
 25 the molecular structures and result desired, according to

what is described in detail in specific literature available to experts in the field.

Examples of these methods are qualitatively schematized in the list provided hereunder, which however does not limit the overall scope of the present invention:

- (m₁) by contact of a metallocene having general formula (X), or preferably (XI), wherein at least one, preferably all, of the substituents Z' and Z'' is hydrogen or an alkyl radical, with an ionic compound whose cation is capable of reacting with one of the substituents to form a neutral compound, and whose anion is voluminous, non-coordinating and capable of delocalizing the negative charge;
- (m₂) by the reaction of a metallocene having the previous formula (X), or preferably (XI), with an alkylating agent, preferably an aluminum trialkyl, used in molar excess of 10/1 to 500/1, followed by the reaction with a strong Lewis acid, such as for example, tris(pentafluorophenyl)borane in more or less stoichiometric quantity or in slight excess with respect to the metal M'';
- (m₃) by contact and reaction of a metallocene having the previous formula (X), or preferably (XI), with a molar excess of 10/1 to 1000/1, preferably from 100/1 to 500/1 of an aluminum trialkyl or an alkylaluminum

halide represented by the following formula



wherein: R^9 is a linear or branched C_1 - C_8 alkyl group,
or one of their mixtures,

5 X is a halogen, preferably chlorine or bromine, and "m" is a decimal number ranging from 1 to 3 extremes included;

10 followed by the addition to the composition thus obtained, of at least an ionic compound of the type described above in such quantities that the ratio between B or Al and the atom M^v in the metallocene complex is within the range of 0.1 to 15, preferably from 1 to 6.

Examples of ionizing ionic compounds or multicomponent
15 reactive systems capable of producing an ionic catalytic system by reaction with a metallocene complex, according to the present invention, are described in the following patent publications, whose content is herein incorporated as reference:

- 20 • European patent application, published under the Nr.: EP-A 277,003, EP-A 277,004, EP-A 522,581, EP-A 495,375, EP-A 520,732, EP-A 478,913, EP-A 468,651, EP-A 427,697, EP-A 421,659, EP-A 418,044;
- 25 • International patent applications published under the Nr.: WO 92/00333, WO 92/05208; WO 91/09882;

- Patents U.S. 5,064,802, U.S. 2,827,446, U.S. 5,066,739.

Non-limiting examples of complex-cocatalyst combinations suitable for the preparation of ionic catalytic systems in accordance with the present invention are schematized hereunder in table (I), with reference to the respective precursors from whose combination they can be obtained. Any compound of any column can be combined, if necessary, with any compound of the remaining ones, according to the method indicated.

Table (I): preparation of ionic catalysts according to the present invention

Method	Metallocene with formula (XI)	Cocatalyst (ii)
5	(THAZ) ₂ ZrMe ₂ (THAZ) ₂ TiMe ₂ (DHBI) ₂ ZrMe ₂ (DHBI) ₂ HfH ₂	[Ph ₃ C] ⁺ [B(C ₆ F ₅) ₄] ⁻
10	(m ₁) (DHBI) CpZrMe ₂ (THAZ) ₂ TiPr ⁱ ₂ (DHBI) ₂ ZrH ₂ (DHBI) ₂ TiBz ₂	[Bu ₃ NH] ⁺ [B(C ₆ F ₅) ₄] ⁻ [PhNMe ₂ H] ⁺ [B(C ₆ F ₅) ₄] ⁻
15	(7,9-Me ₂ -DHBI) ₂ ZrCl ₂ (DHBI) ₂ TiBr ₂ (THAZ) ₂ ZrCl ₂ (DHBI) ₂ Zr(NMe ₂) ₂ (DHBI) ₂ ZrCl ₂	[Ph ₃ C] ⁺ [B(C ₆ F ₅) ₄] ⁻
20	(m ₃) (DHBI) ₂ HfCl ₂ (DHBI) ₂ TiCl ₂ (DHBI)(Ind)ZrCl ₂ (THAZ) ₂ HfCl ₂ (THAZ) ₂ Ti(NMe ₂) ₂	AlEt ₃ [PhNMe ₂ H] ⁺ [B(C ₆ F ₅) ₄] ⁻ AlBu ⁱ ₃ [Bu ₃ NH] ⁺ [B(C ₆ F ₅) ₄] ⁻
25	Abbreviations: Me = methyl, Et = ethyl, Bu = n-butyl, Bu ⁱ = iso-butyl, Ph = phenyl, Bz = benzyl, Pr ⁱ = isopropyl, Cp = cyclopentadienyl Ind = indenyl, DHBI = 4,5-dihydro-benzo[d]indenyl 7,9-Me ₂ -DHBI = 7,9-dimethyl-4,5-dihydro-benzo[d]indenyl , THAZ = 4,5,6-trihydro benzo[e]azulenyl.	

Also included in the scope of the present invention are those catalysts comprising two or more complexes having formula (X) or (XI) mixed with each other. Catalysts of the present invention based on mixtures of complexes having
5 different catalytic activities can be advantageously used in polymerization when a wider molecular weight distribution of the polyolefins thus produced is desired.

In a particularly preferred embodiment of the present invention, the metallocene component (i) of said catalyst
10 consists of a complex selected from those represented by the previous formula (XI), especially with the metal M^a in oxidation state +4, or a mixture of these complexes.

According to another aspect of the present invention, in order to produce solid components for the formation of
15 catalysts for the polymerization of olefins, the above complexes can also be supported on inert solids, preferably consisting of oxides of Si and/or Al, such as, for example, silica, alumina or silica-aluminates. For supporting of said catalysts, the known supporting techniques can be
20 used, normally comprising contact, in a suitable inert liquid medium, between the carrier, optionally activated by heating to temperatures exceeding 200°C, and one or both of components (i) and (ii) of the catalyst of the present invention. For the purposes of the present invention, it is
25 not necessary for both components to be supported, as it is

also possible for only the complex having formula (X), or preferably (XI), or the organic compound of B, Al, Ga or Sn as defined above, to be present on the surface of the carrier. In the latter case, the component which is not present on the surface is subsequently put in contact with the supported component, at the moment of the formation of the catalyst active for the polymerization.

Also included in the scope of the present invention are the complexes, and catalytic systems based on these, which have been supported on a solid by means of the functionalization of the latter and formation of a covalent bond between the solid and a metallocene complex included in formula (X), or preferably (XI), above.

A particular method for the formation of a supported catalyst according to the present invention comprises pre-polymerizing a relatively small fraction of monomer or mixture of monomers in the presence of the catalyst, so as to include this in a solid micro-particulate, which is then fed to the actual reactor itself for completing the process in the presence of an additional α -olefin. This provides a better control of the morphology and dimensions of the polymeric particulate obtained at the end.

One or more other additives or components can be optionally added to the catalyst according to the present invention, as well as the two components (i) or (ii), to ob-

tain a catalytic system suitable for satisfying specific requests. The catalytic systems thus obtained should be considered as being included in the scope of the present invention. Additives or components which can be included in the preparation and/or formulation of the catalyst of the present invention are inert solvents such as, for example, aliphatic and/or aromatic hydrocarbons, aliphatic and aromatic ethers, weakly coordinating additives (Lewis bases) selected, for example, from non-polymerizable olefins, ethers, tertiary amines and alcohols, halogenating agents such as silicon halides, halogenated hydrocarbons, preferably chlorinated, and the like, and again all other possible components normally used in the art for the preparation of the traditional homogeneous catalysts of the metallocene type for the (co)polymerization of ethylene and α -olefins.

Components (i) and (ii) form the catalyst of the present invention by contact with each other, preferably at temperatures ranging from room temperature to 60°C and for times varying from 10 seconds to 1 hour, more preferably from 30 seconds to 10 minutes.

The catalysts according to the present invention can be used with excellent results in substantially all known (co)polymerization processes of α -olefins, either in continuous or batchwise, in one or more steps, such as, for example, processes at low (0.1-1.0 MPa), medium (1.0-10

MPa) or high (10-150 MPa) pressure, at temperatures ranging from 20° to 240°C, optionally in the presence of an inert diluent. Hydrogen can be conveniently used as molecular weight regulator.

5 A further object of the present invention therefore relates to a process for the (co)polymerization of α -olefins, comprising polymerizing said α -olefin, or copolymerizing one or more of said α -olefins, under suitable pressure and temperature conditions, in the presence of at
10 least one of the catalysts according to the present invention, as described above.

These processes can be carried out in solution or suspension in a liquid diluent normally consisting of an aliphatic or cycloaliphatic saturated hydrocarbon, having from
15 3 to 8 carbon atoms, but which can also consist of a monomer as, for example, in the known co-polymerization process of ethylene and propylene in liquid propylene. The quantity of catalyst introduced into the polymerization mixture is preferably selected so that the concentration of the transition metal M^V or M^3 ranges from 10^{-5} to 10^{-6} moles/liter.
20

Alternatively, the polymerization can be carried out in gas phase, for example, in a fluid bed reactor, normally at pressures ranging from 0.5 to 5 MPa and at temperatures ranging from 50 to 150°C.

25 α -olefins which can be polymerized or copolymerized

according to the process of the present invention are generally all olefinically unsaturated hydrocarbons having at least one $>C=CH_2$ double bond in so-called α , or primary, position. Said α -olefins comprise ethylene and its higher homologous products, preferably having from 3 to 15 carbon atoms, such as, for example, propylene, 1-butene, butadiene, 1-hexene, 1-octene, 1,4-hexadiene and styrene. Particular non-primary olefins, having the double bond included in a stretched ring, such as for example, dicyclopentadiene, can be equally (co)polymerized in the presence of the catalyst of the present invention.

According to a particular aspect of the present invention, the catalyst for the (co)polymerization of α -olefins is prepared separately (preformed) by contact of components (i) and (ii), and is subsequently introduced into the polymerization environment. The catalyst can be charged first into the polymerization reactor, followed by the reagent mixture containing the olefin or mixture of olefins to be polymerized, or the catalyst can be charged into the reactor already containing the reagent mixture, or finally, the reagent mixture and the catalyst can be contemporaneously fed into the reactor.

According to another aspect of the present invention, the catalyst is formed "in situ", for example by introducing components (i) and (ii) separately into the polymeriza-

tion reactor containing the selected olefinic monomers.

The catalysts according to the present invention can be used with excellent results in the polymerization of ethylene to give linear polyethylene and in the copolymeri-
5 zation of ethylene with propylene or higher α -olefins, preferably having from 4 to 10 carbon atoms, to give copolymers having different characteristics depending on the specific polymerization conditions and on the quantity and structure of the α -olefin. For example, linear polyethyl-
10 enes can be obtained, with a density ranging from 0.880 to 0.940, and with molecular weights ranging from 10,000 to 2,000,000. The α -olefins preferably used as comonomers of ethylene in the production of low or medium density linear polyethylene (known with the abbreviations ULDPE, VLDPE and
15 LLDPE depending on the density), are 1-butene, 1-hexene and 1-octene.

The catalyst of the present invention can also be conveniently used in copolymerization processes of ethylene and propylene to give saturated elastomeric copolymers vul-
20 canizable by means of peroxides and extremely resistant to aging and degradation, or in the terpolymerization of ethylene, propylene and a non-conjugated diene, having from 5 to 20 carbon atoms, to obtain vulcanizable rubbers of the EPDM type. In the case of these latter processes, it has
25 been found that the catalysts of the present invention al-

low the production of polymers having a particularly high diene content and average molecular weight, under the polymerization conditions.

In the case of the preparation of EPDM, the dienes which can be used for the preparation of said terpolymers are preferably selected from:

- linear chain dienes such as 1,4-hexadiene and 1,6-octadiene;
- branched dienes such as 5-methyl-1,4-hexadiene; 3,7-dimethyl-1,6-octadiene; 3,7-dimethyl-1,7-octadiene;
- dienes with a single ring such as 1,4-cyclohexadiene; 1,5-cyclo-octadiene; 1,5-cyclododecadiene;
- dienes having bridged condensed rings such as dicyclopentadiene; bicyclo[2.2.1]hepta-2,5-diene; alkenyl, alkyldiene, cycloalkenyl and cycloalkyldiene norbornenes such as 5-methylene-2-norbornene, 5-ethylidene-2-norbornene (ENB), 5-propenyl-2-norbornene.

Among the non-conjugated dienes typically used for preparing these copolymers, dienes containing at least one double bond in a stretched ring are preferred, even more preferably 5-ethylidene-2-norbornene (ENB), and also 1,4-hexadiene and 1,6-octadiene.

In the case of EPDM terpolymers, it is convenient for the quantity of dienic monomer not to exceed 15% by weight, and it is preferably from 2 to 10% by weight. The propylene

content on the other hand ranges from 20 to 50% by weight.

The catalysts of the present invention can also be used in homo- and co-polymerization processes of α -olefins according to the known techniques, giving, with excellent yields, atactic, isotactic or syndiotactic polymers, depending on the structure and geometry of the metallocene complex having formula (XI). α -olefins suitable for the purpose are those having from 3 to 20 carbon atoms, optionally also comprising halogens or aromatic nuclei such as, for example, propylene, 1-butene, 1-hexene, 4-methyl-1-pentene, 1-decene and styrene.

The present invention is further described by the following examples, which, however, are provided for purely illustrative purposes and do not limit the overall scope of the invention itself.

EXAMPLES

The analytical techniques and characterization methods used in the examples are listed below and are briefly described.

The characterization by means of ^1H -NMR and ^{13}C -NMR spectroscopy mentioned in the following examples was carried out on a nuclear magnetic resonance spectrometer mod. Bruker MSL-300.

The characterization of the products and organic intermediates by means of gaschromatography/mass spectrogra-

phy (GC-mass) was carried out with a Finnigan TSQ 700 instrument.

The measurement of the molecular weights of the olefinic polymers was carried out by means of Gel-Permeation Chromatography (GPC). The analyses of the samples were effected in 1,2,4-trichlorobenzene (stabilized with Santonox) at 135°C with a WATERS 150-CV chromatograph using a Waters differential refractometer as detector.

The chromatographic separation was obtained with a set of μ -Styragel HT columns (Waters) of which three with pore dimensions of 10^3 , 10^4 , 10^5 Å respectively, and two with pore dimensions of 10^6 Å, establishing a flow-rate of the eluant of 1 ml/min.

The data were obtained and processed by means of Maxima 820 software version 3.30 (Millipore); the number (M_n) and weight (M_w) average molecular weight calculation was carried out by universal calibration, selecting polystyrene standards with molecular weights within the range of 6,500,000-2,000, for the calibration.

The determination of the content of units deriving from propylene and possible diene in the polymers is carried out (according to the method of the Applicant) by means of IR on the same polymers in the form of films having a thickness of 0.2 mm, using an FTIR Perkin-Elmer spectrophotometer model 1760. The intensity of the characteris-

tic peaks is measured, of propylene at 4390 cm^{-1} and ENB at 1688 cm^{-1} respectively, in relation to the peak at 4255 cm^{-1} , and the quantity is determined using a standard calibration curve.

- 5 The Melt Flow Index (MFI) of the polymers is determined in accordance with the regulation ASTM D-1238 D.

In the preparation of the examples, the commercial reagents listed below, were used:

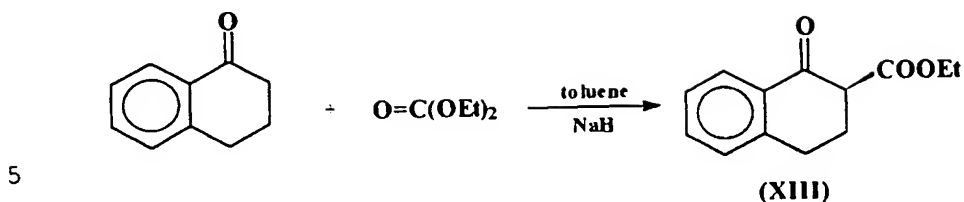
	α -tetralone	ALDRICH
10	α -benzosuberone	ALDRICH
	5,7-dimethyl-1-tetralone	ALDRICH
	diethylcarbonate	CARLO ERBA
	propargylchloride	FLUKA
	methyllithium (LiMe) 1.6 M in diethyl ether	ALDRICH
15	butyllithium (LiBu) 2.5 M in hexane	ALDRICH
	zirconium tetrachloride (ZrCl_4)	FLUKA
	methylalumoxane (MAO) (Eurecene 5100 10T, 10% weight/volume of Al in toluene)	WITCO

- 20 The reagents and/or solvents used and not indicated above are those commonly used both in bench and on industrial scale and can be easily found at all commercial operators specialized in the field.

EXAMPLE 1: Synthesis of bis-(4,5-dihydro-benzo[d]indenyl)-zirconium dichloride (XVIII)

- 25 1a) Synthesis of 1-oxo-2-ethyloxycarbonyl-1,2,3,4-tetra-

hydronaphthalene (XIII)



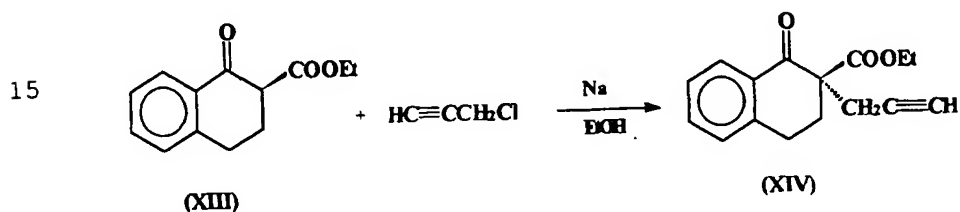
2.5 liters of commercial toluene and 22 g of NaH (0.96 moles) are charged into a 3 liter three-necked glass flask, equipped with a mechanical stirrer and cooler, the suspension is left under stirring at room temperature for 2 hours and 115 ml of diethylcarbonate (0.94 moles) are then added. The mixture is then heated to 80-90°C and 38 ml of α -tetralone (0.28 moles) diluted in 80 ml of toluene are added, in about 3 hours, by means of a drip funnel. The reaction mixture is maintained at 90°C and the trend is periodically controlled by means of thin layer chromatography (T.L.C.) on samples removed for the purpose. After about 24 hours, the complete disappearance of α -tetralone is observed.

20 The reaction mixture is carefully poured into about 1 kg of ice containing 50 ml of glacial acetic acid. The phases are separated and the aqueous phase is repeatedly extracted with ethyl ether, the organic extracts are subsequently washed with water saturated with NaHCO_3 and then
25 with water until neutrality. The organic phase, after anhy-

drification on Na_2SO_4 is evaporated at reduced pressure until 53 g of a red-colored oily residue is obtained, which is purified by distillation on a 20 cm. vigreux column. 42.65 g (density = 1.15 g/cm^3) of the desired 1-oxo-2-ethoxycarbonyl-1,2,3,4-tetrahydronaphthalene having formula (XIII) in the above scheme, are obtained. The NMR spectrum shows that the product is present in solution in ketonic and enolic forms in a ratio of 55/45 respectively.

^1H NMR (CDCl_3 , δ , ppm rel. to TMS): 12.48 (s, 0.45H enol); 7.81-7.14 (m, 4H); 4.24 (dq, 2H); 3.59 (dd, 0.55H ketone); 3.1-2.2 (m, 4H); 1.31 (dt, 3H).

1b) Synthesis of 1-oxo-2-prop-2-ynyl-2-ethoxycarbonyl-1,2,3,4-tetrahydro-naphthalene (XIV)



4.1 g (0.178 moles) of metal sodium are charged into a 1 liter flask and 500 ml of anhydrous ethanol are added. After complete dissolution of the metal sodium, 42 g (0.175 moles) of 1-oxo-2-ethoxycarbonyl-1,2,3,4-tetrahydronaphthalene (XIII) obtained as described above, are added. A light yellow milky solid is immediately formed. The mixture is heated to the reflux temperature of the solvent, and 42 ml (0.4 moles) of propargyl chloride diluted with 30% of

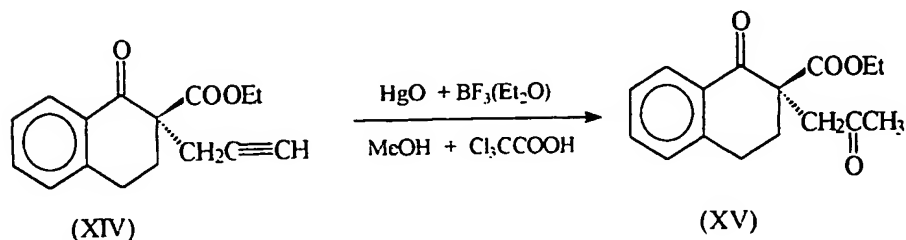
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toluene are then added by means of a drip funnel in about 2 hours. At the end of the addition, the mixture is kept at reflux temperature for about 3 hours and is then cooled to room temperature and left under stirring for the whole
5 night. In this phase there is the progressive disappearance of the yellow milky solid with the formation of a colorless crystalline solid. About 400 ml of EtOH are removed by evaporation at reduced pressure, the mixture is subsequently hydrolyzed with water and ice and extracted with
10 various portions of ethyl ether. The ether extracts are joined, washed with water until neutrality and dried on anhydrous Na_2SO_4 . The solvent is then evaporated at reduced pressure and 45 g of an oily reddish-brown residue is obtained, consisting of 70% by weight of the desired product
15 (XIV) (GC analysis, 70% yield).

^1H NMR (CDCl_3 ; δ , ppm rel. to TMS): 7.4-7.1 (m, 4H); 4.14 (q, 2H); 3.2-2.95 (m, 2H); 2.88 (d, 2H); 2.7-2.3 (m, 2H); 2.00 (t, 1H); 1.53 (t, 3H).

1c) Synthesis of 1-oxo-2-(2-oxo-propyl)-2-ethoxycarbonyl-
20 1,2,3,4-tetrahydro-naphthalene (XV):



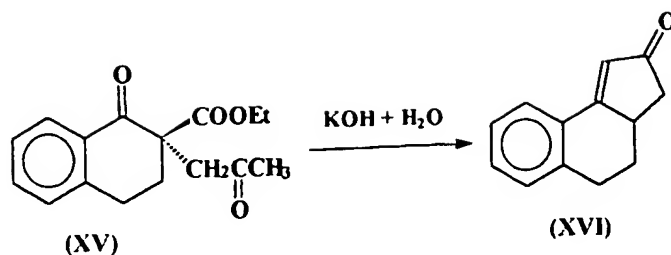
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The following products are charged in order into a 500 ml glass flask: 200 ml of methanol, 0.32 g (1.5 mmols) of mercury oxide, 1.4 ml (11 mmols) of borontrifluoride etherate and 0.07 g (0.4 mmols) of trichloroacetic acid.

5 45 g (0.123 moles) of the compound having formula (XIV) obtained as described in the previous step (b), dissolved in 50 ml of methanol, are added to the suspension thus obtained. The reaction mixture is then left under stirring at room temperature for 18 hours and is subsequently hydrolyzed with water and extracted with various
10 portions of ethyl ether. The ether extracts are joined, anhydri-fied on Na_2SO_4 and concentrated by evaporation at reduced pressure, obtaining at the end an oily red residue containing 43.5 g of the desired product having formula
15 (XV) (65% purity upon GC analysis, 86% yield). Analyses show that both ethyl ester (45%) and methyl ester (55%) deriving from the transesterification of the product (XV) due to the use of methanol as solvent, are present. The product is not additionally purified.

20 ^1H NMR (CDCl_3 , δ , ppm rel. to TMS): 7.4-7.1 (m, 4H); 4.14 (q, 0.45x2H, ethyl ester); 3.67 (s, 0.55x3H, methyl ester); 3.2-2.3 (m, 6H); 2.22 (s, 3H); 1.29 (t, 0.45x3H, ethyl ester).

25 1d) Synthesis of 3,3a,4,5-tetrahydro-cyclopenta[a]naphthalen-2-one (XVI)



5

500 ml of water and 40 g of KOH are charged into a 1 liter glass flask, containing 43 g of 2-ethoxycarbonyl-2-(2-oxopropyl)- α -tetralone (XV), 65% purity, obtained as described above. The mixture is then heated to reflux temperature for 10 hours, the reaction trend being followed by means of TLC.

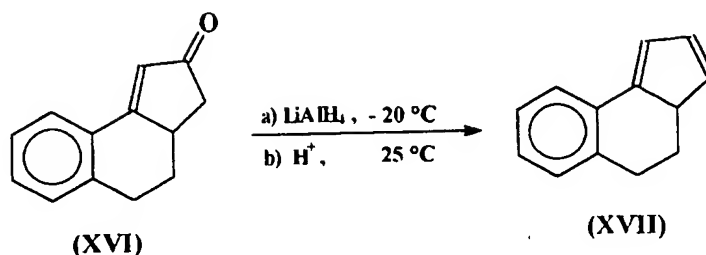
The aqueous suspension is cooled to room temperature and extracted with various portions of ethyl ether, the organic fractions are joined and subsequently washed with water saturated with NH_4Cl , then with water until neutrality and dried on anhydrous Na_2SO_4 . The solvent is removed under vacuum and 26.5 g of an oily red residue are obtained, which contains 65% (GC analysis) of the product (XVII) (yield 93%), which is purified by means of flash chromatography, using a 70/30 (by volume) mixture of hexane/ethyl acetate as eluant and SiO_2 (Merck) 32-60 Å as stationary phase. At the end of this operation, 9.4 g (51 mmols) of the desired pure product having formula (XVII) are recovered.

^1H NMR (CDCl_3 ; δ , ppm rel. to TMS): 7.65 (d, 1H); 7.4-7.2

(m, 3H); 6.38 (d, 1H,); 3.1-2.9 (m, 3H); 2.84-2.72 (dd, 1H); 2.35-2.26 (m, 1H); 2.25-2.14 (dd, 1H); 1.77-1.55 (m, 1H).

le) Synthesis of 3a,4,5-trihydro-cyclopenta[a]naphthalene

5 (XVII)

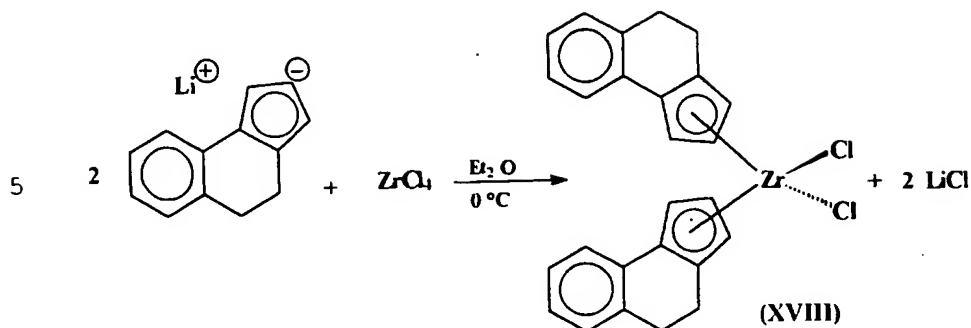


10

3.5 g (92 mmoles) of LiAlH_4 and 250 ml of ethyl ether are charged into a 500 ml flask. 50 ml of a solution of ethyl ether containing 4.0 g (21,5 mmoles) of the product having formula (XVI) are added at -20°C by means of a drip
15 funnel over a period of about 1 hour, to the suspension thus formed. At the end of the addition, the temperature is brought to 25°C and the mixture is left under stirring for 2 hours, obtaining a yellow mixture. This reaction mixture is carefully poured into water and ice and is then acidified
20 with HCl , extracted with various portions of ethyl ether, the organic phase is dried on anhydrous Na_2SO_4 , and finally the solvent is evaporated at reduced pressure, obtaining 2.9 g of the desired pure product (XVII), as a light yellow granular solid (yield 80%).

25 lf) Synthesis of bis-(4,5-dihydro cyclopenta[a]naphthal-

enyl zirconium dichloride (XVIII).



2.9 g (17.3 mmol) of the cyclopentadienyl compound
 10 having formula (XVII) obtained according to the previous
 step (e) and 200 ml of hexane are charged into a 500 ml
 flask. 16 ml (25.6 mmol) of LiBu (1.6 M in hexane) are
 then added in about 30 minutes by means of a drip funnel. A
 yellow solid is rapidly formed. The mixture is left under
 15 stirring at room temperature for 18 hours, the solid is
 then recovered by filtration, washing with three 20 ml por-
 tions of hexane and the product is dried at reduced pres-
 sure, thus obtaining 0.7 g of the lithium salt of the cy-
 clopentadienyl compound having formula (XVII) (24% yield).

20 The lithium salt thus obtained and 100 ml of ethyl
 ether are charged into a 250 ml flask. A suspension is ob-
 tained which is cooled to 0°C and 0.5 g (2.14 mmol) of
 ZrCl₄ are added. The reaction mixture is then left under
 stirring for 4 hours in this phase; a yellowish colouring
 25 is obtained and a white solid is formed which can be easily

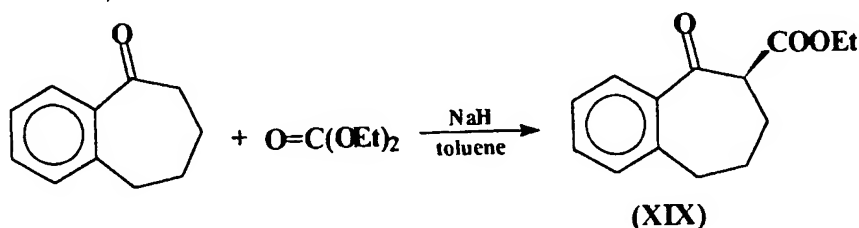
decanted. The suspension is then filtered and the solid washed with three 10 ml portions of ethyl ether. The ether solution is evaporated under vacuum, at room temperature, and in this way a solid yellow residue remains, from which
5 the complex (XVIII), pure by crystallization from toluene pentane, is obtained.

¹H NMR (CD₂Cl₂, δ, ppm rel. to TMS): 7.41 (d, 1H); 7.35-7.15 (m, 7H); 6.36 (t, 1H); 6.30 (dt, 2H); 6.05 (dd, 1H); 5.90 (dd, 1H); 5.77 (t, 1H); 3.10-2.95 (m, 3H); 2.90-2.60 (m,
10 9H).

EXAMPLE 2: Synthesis of bis-(4,5,6-trihydro-benzo[e]azul-enyl)zirconium dichloride (XXIV)

2a) Synthesis of 5-oxo-6-ethoxycarbonyl-6,7,8,9-tetrahydro-5H-benzocycloheptadiene (XIX).

15

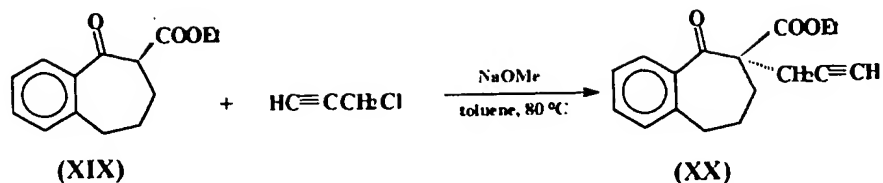


20 11.5 g of NaH (0.48 moles) in 600 ml of commercial toluene are suspended in a 1 liter glass flask, equipped with a bubble cooler, and the whole mixture is maintained under stirring at room temperature for 2 hours. 63 ml of diethyl carbonate (0.52 moles) are then added and the mixture is heated to 90°C, after which 25 g of α-benzosuberone
25

(0.156 moles) dissolved in 50 ml of toluene are added, by means of a drip funnel in about 3 hours. The reaction trend is controlled by means of T.L.C. and after about 4 hours at 90°C, the complete disappearance of α -benzosuberone is observed. The reaction mixture is carefully poured into about 1 kg of ice containing 50 ml of glacial acetic acid. The phases are separated and the aqueous phase is repeatedly extracted with ethyl ether, the organic extracts are subsequently joined and washed with water saturated with NaHCO₃ and then with water until neutrality. The organic phase is anhydriified on Na₂SO₄ and the solvents are completely removed at reduced pressure. 33.2 g of a yellow oil consisting of the desired ketoester having formula (XIX) in the above scheme, are obtained, with a 90% purity. The ¹H NMR spectrum shows that the product (XIX) is present in solution in ketonic and enolic forms in a ratio of 25 to 75% respectively.

¹H NMR (CDCl₃, δ , ppm rel. to TMS): 12.70 (s, 0.75H enol); 7.7-7.1 (m, 4H); 4.28 (dq, 2H); 3.78 (dd, 0.25H ketone); 2.93 (m, 2x0.25H ketone); 2.63 (t, 2x0.75H enol); 2.3-1.8 (m, 4H); 1.34 (t, 3x0.75H enol); 1.25 (t, 3x0.25H ketone).

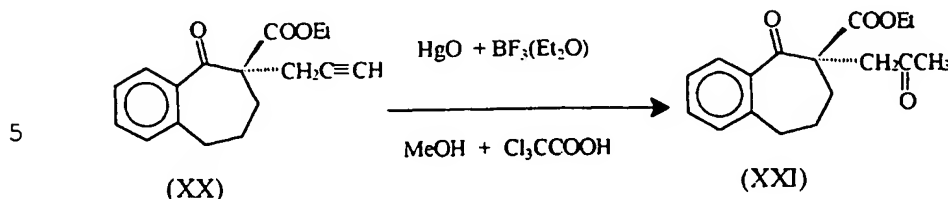
2b) Synthesis of 5-oxo-6-ethoxycarbonyl-6-prop-2-ynyl-6,7,8,9-tetrahydro-5H-benzocycloheptadiene (XX).



5 4.5 g of metal sodium and 100 ml of anhydrous methanol are charged into a 500 ml glass flask. The mixture is left under stirring until complete dissolution of the sodium, after which the solvent is evaporated and the solid obtained is further dried in order to remove most of the
10 methanol present.

The colourless solid thus obtained is dispersed in 200 ml of toluene, the ketoester (XIX) is added, the suspension is then heated to 80°C and 31 ml (0.295 moles) of propargyl chloride in a solution at 70% in toluene are added, by
15 means of a drip funnel, in about an hour. After 24 hours, G.C. analysis of the reaction mixture shows that all the ketoester (XIX) has been converted. The reaction mixture is then hydrolyzed with water and ice and extracted with various portions of ethyl ether. The ether extracts are
20 joined and subsequently washed with water and dried on anhydrous Na₂SO₄. The solvent is then removed at reduced pressure and 38.5 g of a reddish-brown oil are thus obtained, in which about 35% of the desired product having formula (XX) is present, as determined on the basis of GC-
25 mass analysis.

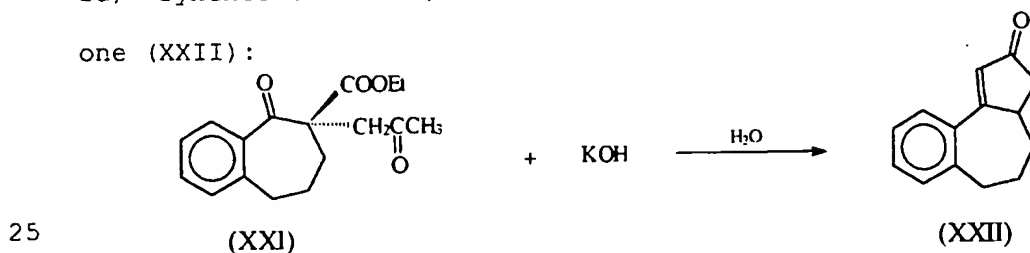
2c) Synthesis of 5-oxo-6-ethoxycarbonyl-6-(2-oxo-propyl)-6,7,8,9-tetrahydro-5H-benzocycloheptadiene (XXI):



The following products are charged in order into a 500 ml flask, at room temperature: 250 ml of methanol, 1.2 g (5.6 mmoles) of mercury oxide HgO , 1.5 ml (11.8 mmoles) of boron trifluoride etherate, 0.07 g (0.4 mmoles) of trichloroacetic acid and 38.5 g of the impure product (35%) having formula (XX) obtained as described above. The dark suspension thus formed is reacted by heating to 50°C for 4 hours. At the end, the methanol is removed at reduced pressure and water is added to the residue, forming a suspension which is extracted with various portions of ethyl ether. These are joined, washed with water until neutrality and anhydri-
fied on Na_2SO_4 . The ether solution is dried obtaining 38 g of a reddish-brown oil containing about 35% of the desired product (XXI).

15
20

2d) Synthesis of 3,4,5,6-tetrahydro-3H-benzo[e]azulen-2-one (XXII):

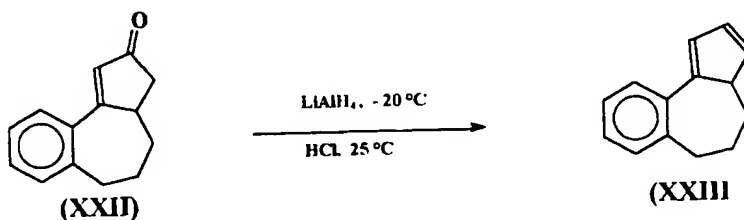


55 g (0.98 moles) of potassium hydroxide are dissolved in 300 ml of water in a 500 ml flask and 38 g of compound (XXI), obtained as described above, are added. A suspension is formed, which is heated to reflux temperature for 12 hours, the reaction being followed by G.C.

At the end of the reaction, the suspension is cooled to room temperature and extracted with various portions of ethyl ether. The ether extracts are subsequently washed with water saturated with NH_4Cl and subsequently with water until neutrality, then dried on anhydrous Na_2SO_4 . After evaporation of the solvent at reduced pressure, 13 g of an oily red liquid are obtained, which is purified by means of flash chromatography on a column, using a 70/30 (by volume) mixture of hexane/ethyl acetate as eluant and SiO_2 (Merck) 32-60 as stationary phase. 2.2 g of the desired pure product (XXII) are obtained at the end.

^1H NMR (CDCl_3 , δ , ppm rel. to TMS): 7.4-7.1 (m, 4H); 6.20 (d, 1H); 3.2-2.6 (m, 4H); 2.4-1.2 (m, 5H).

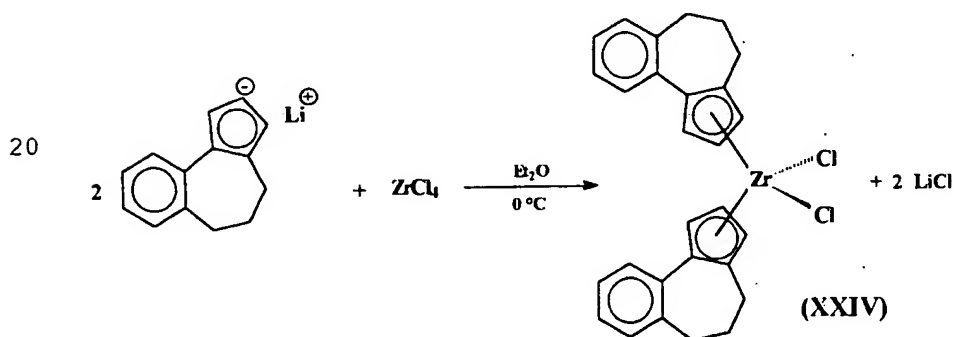
2e) Synthesis of 3a,4,5,6-tetrahydro-benzo[e]azulene (XXIII)



25

A solution of 2.2 g of the compound (XXII), prepared as described above, in 20 ml of ethyl ether, are charged, over a period of about 1 hour, into a 250 ml flask cooled to -20°C , containing a mixture of 2.1 g (55 mmols) of LiAlH_4 and 100 ml of ethyl ether. A light yellow suspension is formed which is maintained under stirring at room temperature for a further 2 hours. At the end of the reaction, the suspension is carefully poured into water and ice and is then acidified with HCl and extracted with four 50 ml portions of ethyl ether. The organic phase is treated with p-toluenesulfonic acid in excess and maintained at 4°C for 18 hours. It is then hydrolyzed with water saturated with NaHCO_3 and washed until neutrality. After drying on Na_2SO_4 and removing the solvent under vacuum, 1.1 g of the desired product (XXIII) are obtained in the form of a red oil.

2f) Synthesis of bis-(4,5,6-trihydro-benzo[e]azulenyl)-zirconium dichloride (XXIV)



1.1 g (6 mmoles) of the product (XXIII), obtained as described above, are dissolved in ethyl ether in a 100 ml flask and 10 ml of lithium butyl 1.6 M in hexane (16 mmoles) are added, after cooling to -20°C, over a period of about 30 minutes. The mixture is then brought to room temperature and left to react, under stirring, for a further 4 hours. A light yellow solid is formed, which is recovered by filtration, washed three times with 10 ml of hexane and dried under vacuum.

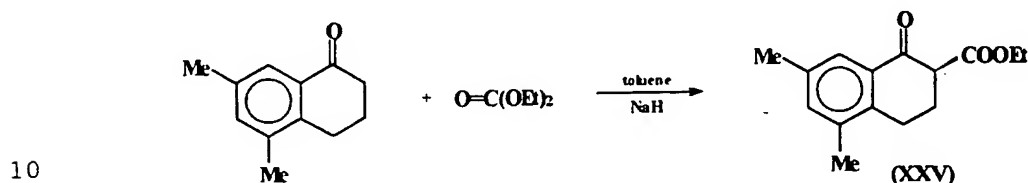
0.94 g of lithium salt of compound (XXIII) are thus obtained (83% yield), which are charged into a 250 ml flask containing 100 ml of ethyl ether. The suspension thus obtained is cooled to 0°C, 0.52 g of ZrCl₄ are added and the mixture is maintained under stirring for 4 hours. The reaction mixture is brought to room temperature, filtered and the solid extracted three times with 10 ml of ethyl ether. After evaporating the ether, an orange oily residue is obtained which is treated under heat with a mixture of toluene/hexane (1/1 by volume). The red insoluble part is removed by filtration under heat and on cooling the hydrocarbon solution, a yellow-orange solid is formed, which is recovered by filtration, washed with hexane (10 ml) and dried under vacuum. 0.21 g of the desired zirconium complex (XXIV) are thus obtained.

¹H NMR (CDCl₃ δ, ppm rel. to TMS): 7.4-7.0 (m, 8H); 6.63

(t, 1H); 6.60 (t, 1H); 6.21 (m, 4H); 3.1-2.9 (m, 2H); 2.8-2.4 (m, 6H); 2.3-1.9 (m, 4H).

EXAMPLE 3: Synthesis of bis-(7,9-di-methyl-4,5-dihydro-benzo[d]indenyl)zirconium dichloride (XXX)

- 5 3a) Synthesis of 5,7-dimethyl-1-oxo-2-ethoxycarbonyl-1,2,3,4-tetrahydronaphthalene (XXV)



A suspension of 12.2 g (0.53 moles) of sodium hydride in a liter of toluene are maintained under stirring in a 2 liter flask for 2 hours at room temperature, and 85 ml (0.69 moles) of diethylcarbonate are then added. The mixture is heated to 90°C and 25 g of 5,7-dimethyl-1-tetralone (0.14 moles) dissolved in 150 ml of toluene are added, in about 2 hours. The reaction mixture is kept under stirring for a further 8 hours at 90°C until the reagent disappears.

20 At the end the reaction mixture is hydrolyzed by pouring it into ice containing 50 ml of glacial acetic acid. The phases are separated and extracted repeatedly with ethyl ether, the overall organic phase is washed with water saturated with NaHCO₃, then with water until neutrality and finally dried on anhydrous Na₂SO₄. Upon removal of the sol-

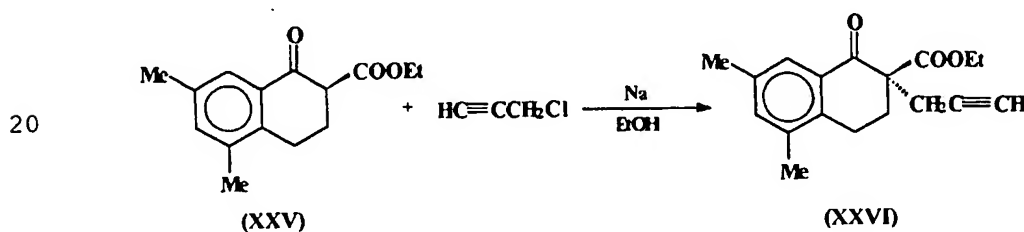
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vents at reduced pressure, 33 g of the desired pure ketoester (XXV) are obtained in the form of a red oil (96% yield). NMR spectra show that the product is present in solution in ketone and enolic form in a ratio of 55% and 45% respectively.

^1H NMR (CDCl_3 ; δ , ppm rel. to TMS): 12.50 (s, 0.45H); 7.71 (s, 0.55H); 7.49 (s, 0.45H); 7.17 (s, 0.55H); 7.01 (s, 0.45H); 4.25 (dq, 2H); 3.53 (dd, 0.55H); 3.1-2.2 (m, 4H); 2.30 (s, 3H); 2.24 (s, 3H); 1.34 (t, 0.45x3H); 1.28 (t, 0.55x3H).

^{13}C NMR (CDCl_3 ; δ , ppm rel. to TMS): 194.38; 173.46; 170.96; 166.18; 136.77; 136.96; 136.63; 136.08; 135.44; 134.01; 132.66; 130.64; 129.81; 128.93; 97.07; 61.79; 61.11; 54.73; 26.43; 25.11; 24.23; 21.64; 21.47; 21.04; 19.89; 19.84; 15.04; 14.87.

3b) Synthesis of 5,7-dimethyl-1-oxo-2-ethoxycarbonyl-2-(prop-2-ynyl)-1,2,3,4-tetrahydronaphthalene (XXVI)



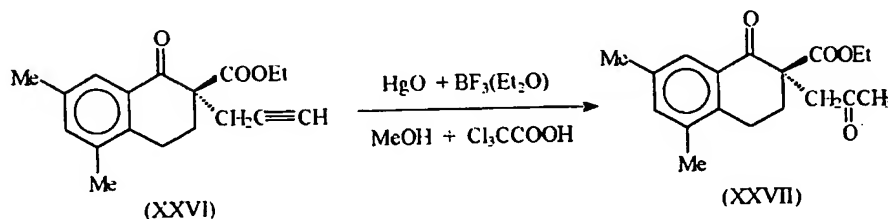
5.0 g (0.0217 moles) of metal sodium and 500 ml of anhydrous ethyl alcohol are charged into a 1 liter flask and kept under stirring until the complete dissolution of the

sodium; the ketoester (XXV) is then added. 30 ml of propargyl chloride (0.285 moles) dissolved in 70 ml of toluene and 20 ml of EtOH are then added, by means of a filter funnel, over a period of 1 hour. The suspension thus obtained is maintained under stirring at reflux temperature for a further 8 hours, is then cooled to room temperature, hydrolyzed by the addition of water and extracted with various portions of ethyl ether. The organic phase is repeatedly washed with water until neutrality and dried on anhydrous Na_2SO_4 . The solvent is subsequently evaporated under vacuum, thus obtaining 25 g of a red oil which, upon subsequent ^1H NMR analysis proves to be the desired product (XXVII) (65% yield).

^1H NMR (CDCl_3 , δ , ppm rel. to TMS): 7.72 (s, 1H); 7.19 (s, 1H); 4.13 (dq, 2H); 3.1-2.6 (m, 2H); 2.86 (dd, 2H); 2.5-2.2 (m, 2H); 2.31 (s, 3H), 2.24 (s, 3H); 2.01 (t, 1H); 1.14 (t, 3H).

3c) Synthesis of 5,7-dimethyl-1-oxo-2-ethoxycarbonyl-2-(2-oxo-propyl)-1,2,3,4-tetrahydronaphthalene (XXVII)

20

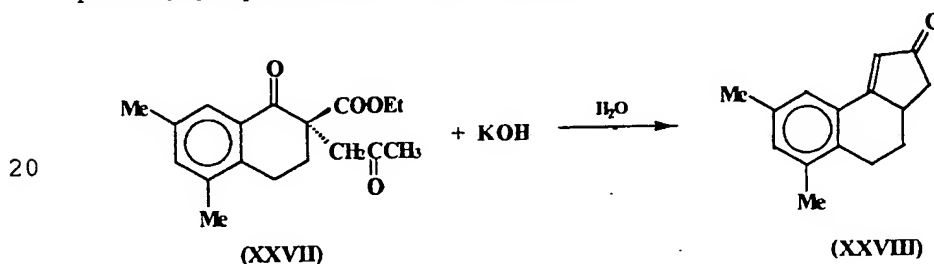


The following products are charged in order into a 1 liter flask: 1.5 g (7 μmoles) of HgO , 2 ml (15.7 μmoles) of

boron trifluoride etherate ($\text{BF}_3 \cdot (\text{Et}_2\text{O})$), 0.015 g (0.9 mmoles) of trichloroacetic acid, 400 ml of methanol and 20 g (0.07 moles) of the product (XXVI) obtained as described above. The reaction mixture is heated to 50°C and maintained at this temperature, under stirring for 5 hours. After cooling to room temperature, water is added to hydrolyze the mixture, which is then extracted with various portions of ethyl ether. 19.3 g of an oily liquid are obtained from the organic phase, after anhydrifying on Na_2SO_4 and removal of the solvent at reduced pressure, which, after characterization, proves to be the desired product having formula (XXVII) (96% yield).

^1H NMR (CDCl_3 , δ , ppm rel. to TMS): 7.71 (s, 1H); 7.18 (s, 1H); 3.66 (s, 3H); 3.01 (s, 2H); 2.79 (m, 2H); 2.45 (m, 2H); 2.31 (s, 3H); 2.23 (s, 3H); 2.22 (s, 3H).

3d) Synthesis of 6,8-dimethyl-3,3a,4,5-tetrahydro-cyclopenta[a]naphthalen-2-one (XXVIII)



The following products are charged in order into a 500 ml flask: 250 ml of water, 30 g (0.53 moles) of potassium hydroxide and 10 g (34.7 mmoles) of the product having for-

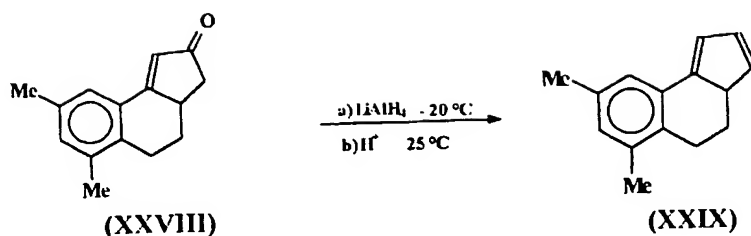
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mula (XXVII) obtained as described above. The mixture is then heated to reflux temperature for 50 hours, the reaction trend being controlled by T.L.C. At the end, a suspension is obtained which is cooled to room temperature and
5 extracted with various portions of ethyl ether, the overall organic phase is then washed with water saturated with NH_4Cl and subsequently with water until neutrality, and is finally dried on anhydrous Na_2SO_4 . After removal of the solvent at reduced pressure, 3.5 g of brown semi-solid
10 residue are obtained, which contains about 70% of the desired cyclopentenone product having formula (XXVIII) (33% yield). The pure product is then recovered by means of flash chromatography, using a 70/30 (by volume) mixture of hexane/ethyl acetate as eluant and SiO_2 (Merck) 32-60 as
15 stationary phase.

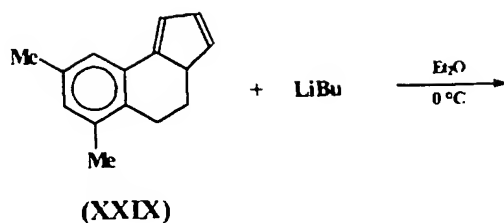
^1H NMR (CDCl_3 ; δ , ppm rel. to TMS): 7.31 (s, 1H); 7.08 (s, 1H); 6.35 (d, 1H); 3.1-2.6 (m, 4H); 2.4-2.1 (m, 2H); 2.32 (s, 3H); 2.24 (s, 3H); 1.8-1.5 (m, 1H).

^{13}C NMR (CDCl_3 ; δ , ppm rel. to TMS): 208.87; 176.97; 137.75;
20 136.42; 135.48; 134.50; 130.55; 125.82; 123.96; 108.24; 107.23; 43.44; 40.09; 30.46; 27.47; 21.53; 20.13.

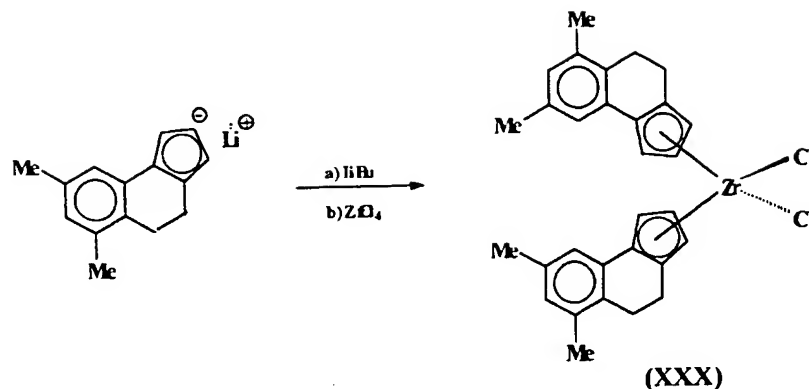
3e) Synthesis of 6,8-dimethyl-3a,4,5-trihydrocyclopenta[a]naphthalene (XXIX)



- 5 1.2 g (31 mmol) of LiAlH_4 are suspended in 50 ml of diethyl ether in a 100 ml flask and, after bringing the temperature to -20°C , a solution of 1.6 g (7.5 mmol) of the product having formula (XXX), obtained as described above, in 20 ml of Et_2O , are added over a period of about
- 10 30 minutes. At the end of the addition the temperature is brought to 25°C and the mixture is maintained under stirring for a further 3 hours. The reaction mixture is then hydrolyzed by carefully pouring it into water and ice, is acidified by adding 5 ml of conc. HCl , extracted with ethyl
- 15 ether and the organic phase is anhydri-fied on Na_2SO_4 . After evaporation of the solvent under vacuum, 1.4 g of a light brown solid product are obtained, which, after characterization, proves to be the desired substituted cyclopentadiene product having formula (XXIX), almost pure.
- 20 3f) Synthesis of bis-(6,8-dimethyl-4,5-dihydrocyclopenta-[a]naphthalenyl)zirconium dichloride (XXX)



5



10

75 ml of ethyl ether and 1.4 g (6.6 mmols) of the diene having formula (XXIX) obtained as described above, are charged into a 250 ml flask. After cooling the mixture to 0°C, 10 ml of a solution of lithium butyl 1.6 M in hexane (16 mmols) are added by means of a syringe, the temperature is left to slowly rise to 25°C and the mixture is maintained under stirring for 18 hours. The solvent is then removed at reduced pressure and hexane (100 ml) is added.

The light brown solid which is formed in this phase is recovered by filtration, washed on a filter with 3 20 ml portions of hexane and dried under vacuum. 0.3 g of the lithium salt of the product having formula (XXIX) are thus obtained.

This lithium salt (1.48 mmols) is suspended in 100 ml

of anhydrous ethyl ether in a 250 ml flask and, after cooling to -20°C , 0.16 g (0.69 mmoles) of ZrCl_4 are added. The temperature is left to rise to 25°C and the mixture is then kept under stirring for 4 hours. A light-coloured solid is
5 formed which can be easily decanted. The suspension is filtered, the resulting ether solution is then dried and the solid thus obtained is re-crystallized from toluene/pentane. The desired metallocene complex having formula (XXX) is thus recovered.

10 EXAMPLES 4-14: copolymerization of ethylene with MAO as co-catalyst.

Examples 4 to 14 refer to a series of co- and terpolymerization tests for the preparation of elastomeric polymers of the EP(D)M type based on ethylene/propylene/ethylidenenorbornene (ENB), carried out using
15 a catalytic system comprising one of the metallocene complexes, obtained as described above in examples 1 and 2, and methylalumoxane (MAO) as cocatalyst. The specific polymerization conditions of each example and the results obtained are indicated in Table (II), which specifies in succession, the reference example number, the metallocene complex used, the quantity of zirconium used, the atomic ratio between aluminum in MAO and zirconium in the metallocene, the total polymerization pressure, the initial concentration of ENB, if present, the activity of the catalytic sys-
20
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tem as kilograms of polymer per gram of metal zirconium per hour ($\text{kg}_{\text{pol.}}/\text{g}_{\text{Zr}}\cdot\text{h}$), the relative quantity, by weight, of the C_2 monomeric units and ENB in the polymer, the weight average molecular weight M_w and the molecular weight dispersion

5 M_w/M_n .

The polymerization is carried out in a 0.5 liter pressure reactor, equipped with a magnetic anchor drag stirrer and an external jacket connected to a heat exchanger for the temperature control. The reactor is previously flushed

10 by maintaining under vacuum (0.1 Pascal) at a temperature of 80°C for at least 2 h.

120 g of liquid "polymerization grade" propylene and the ENB diene, if required, are fed into the reactor at 23°C , in such a quantity as to obtain the molar concentration indicated in the corresponding column in table (II)

15 below. The reactor is then brought to the polymerization temperature of 40°C and, "polymerization grade" gaseous ethylene is fed by means of a plunged pipe until the total desired equilibrium pressure (2.0-2.7 MPa) is reached. Under

20 these conditions the molar concentration of ethylene in the liquid phase ranges from 11 to 23%, depending on the total pressure of the system, as can be easily calculated using the appropriate liquid-vapor tables.

MAO, as a 1.5 M solution (as Al) in toluene and the

25 desired quantity of one of the above metallocene complexes,

as a toluene solution having a concentration generally ranging from 3×10^{-4} to 1×10^{-3} M, are charged into a suitable tailed test-tube, maintained under nitrogen. The catalyst solution thus formed is maintained at room temperature for a few minutes and is then transferred under a stream of inert gas to a metal container from which it is introduced into the reactor, by means of an overpressure of nitrogen.

The polymerization reaction is carried out at 40°C, care being taken that the total pressure is kept constant by continuously feeding ethylene to compensate the part which has reacted in the meantime. After 15 minutes the feeding of ethylene is interrupted and the polymerization is stopped by the rapid degassing of the residual monomers. The polymer is recovered, after washing it with ethyl alcohol and drying at 60°C, at a reduced pressure of 1000 Pa, for at least 8 hours, in order to completely eliminate any possible residual monomers. The solid thus obtained is weighed and the catalytic activity is calculated as described above. The content of the various C_3 monomeric units and ENB is measured on the dried and homogenized solid, by means of the known techniques based on IR spectroscopy, together with the weight (M_w) and number (M_n) average molecular weight. The results are indicated in Table II.

Example Nr.	Complex (formula)	Zr (molesx10 ⁵)	Al/Zr (mol./mol.)	P _{total} (MPa)	ENB _{initial} (% moles)	Activity (kg _{PE} /g _{Zr} ·h)	C ₃ (polymer) (weight%)	ENB (polymer) (weight%)	M _w (x 10 ³)	M _w /M _n
4	(XVIII)	0.40	3950	2.5	---	3360	34.5	---	378	3.1
5	(XVIII)	0.53	3600	2.2	---	912	53.3	---	131	3.0
6	(XVIII)	0.66	2350	2.5	0.4	1299	45.9	3.5	105	2.9
7	(XVIII)	0.27	5850	2.5	---	4721	38.5	---	160	3.8
8	(XVIII)	1.06	2230	2.5	0.8	1120	38.4	3.7	166	2.5
9	(XVIII)	0.20	7900	2.7	---	10769	36.3	----	229	3.0
10	(XVIII)	1.33	2200	2.7	1.2	793	33.4	4.9	218	2.7
11	(XXIV)	0.31	4950	2.5	---	1560	31.0	---	148	2.7
12	(XXIV)	0.43	3580	2.7	---	1533	28.3	---	362	2.9
13	(XXIV)	1.55	1990	2.2	---	1191	37.3	---	243	3.1
14	(XXIV)	0.93	3310	2.4	---	1134	31.8	---	138	2.8

TABLE II: ethylene co- and ter-polymerization according to examples 4 to 14

EXAMPLES 15 TO 19:

Examples 15 to 19 refer to copolymerization tests of ethylene with propylene for the preparation of the corresponding elastomeric polymers of the EPR type, carried out using a catalytic system comprising the metallocene complex having formula (XVIII) obtained as described above in example 1, an aluminum alkyl and an appropriate compound of boron as co-catalyst (catalytic system of the cationic type). The specific polymerization conditions of each example and the results obtained are indicated in Table III, which specifies in succession, the reference example number, the quantity of zirconium used, the atomic ratio between aluminum and zirconium, the atomic ratio between boron and zirconium, the total polymerization pressure, the activity of the catalytic system with reference to the zirconium, the relative quantity, by weight, of the C_3 monomeric units in the polymer, the weight average molecular weight M_w and the molecular weight dispersion M_w/M_n .

Preparation of the catalytic system

$Al(iso-Bu)_3$ as a 0.4 M solution in toluene and the desired quantity of the metallocene complex (XVIII), as a toluene solution having a concentration generally ranging from 3×10^{-4} to 1×10^{-3} M, are charged into a suitable tailed test-tube, maintained under nitrogen. The solution thus formed is maintained under stirring at 23°C for 15 minutes,

after which a toluene solution, having a concentration generally ranging from 5×10^{-4} to 1×10^{-3} M, of $[\text{CPh}_3][\text{B}(\text{C}_6\text{F}_5)_4]$ is added, and then, after a few minutes, is transferred under a stream of inert gas to a metal container from which it is introduced into the reactor, by means of an overpressure of nitrogen.

Polymerization

The polymerization is carried out in a 0.5 liter pressure reactor, equipped with a magnetic anchor drag stirrer and an external jacket connected to a heat exchanger for the temperature control. The reactor is previously flushed by maintaining under vacuum (0.1 Pascal) at a temperature of 80°C for at least 2 h.

120 g of liquid "polymerization grade" propylene and the exact quantity of $\text{Al}(\text{iso-Bu})_3$ so as to obtain an aluminum concentration of 5×10^{-3} moles/liter, are fed into the reactor at 23°C . The reactor is then brought to the polymerization temperature of 40°C and, "polymerization grade" gaseous ethylene is fed by means of a plunged pipe until the desired equilibrium pressure (2.2-2.7 MPa) is reached. Under these conditions the molar concentration of ethylene in the liquid phase ranges from 12 to 23%, depending on the total pressure of the system, as can be easily calculated using the appropriate liquid-vapor tables. At this point the catalytic system is introduced from a suitable con-

tainer connected to the reactor, by means of an overpressure of nitrogen. The polymerization reaction is carried out using the same procedure and under the same conditions as the previous examples 4 to 14. At the end, the polymer
5 thus obtained is weighed and the catalytic activity is calculated as kilograms of polymer per gram of metal zirconium per hour ($\text{kg}_{\text{pol.}}/\text{g}_{\text{Zr}}\cdot\text{h}$). The content of propylene units is measured on the dried and homogenized solid, by means of the known techniques based on IR spectroscopy, together
10 with the weight (M_w) and number (M_n) average molecular weight. The results are indicated in Table (III).

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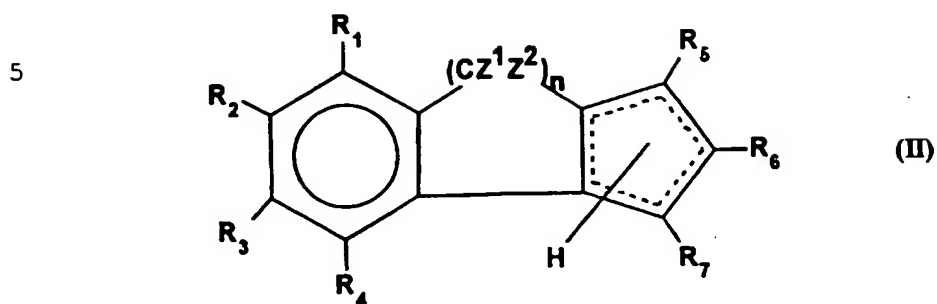
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Example Nr.	Zr (moles $\times 10^6$)	Al/Zr (mol/mol)	B/Zr (mol/mol)	P _{total} (MPa)	Activity (kg _{pol} /g _{Zr} ·xh)	C ₃ (polymer) (weight %)	M _w ($\times 10^3$)	M _w /M _n
15	0.38	316	1.1	2.2	1157	40.4	180	3.4
16	0.38	316	1.1	2.5	1619	35.5	281	3.3
17	0.38	316	4.3	2.2	1215	44.3	186	2.5
18	0.38	316	2.1	2.5	1677	36.3	215	2.9
19	0.38	316	4.3	2.5	1735	37.2	214	2.6

TABLE III : Ethylene co-polymerization according to examples 15 to 19

CLAIMS

1. A polycyclic cyclopentadienyl compound having the following formula (II):



- 10 wherein: each of the symbols R_1 , R_2 , R_3 , R_4 , R_5 , R_6 , R_7 , Z^1 and Z^2 independently represents hydrogen or an organic substituent having from 1 to 15 carbon atoms, and in addition, any one of said R or Z groups can be a divalent organic group further
- 15 bound to another organic group having from 5 to 20 carbon atoms and comprising a cyclopentadienyl group, and

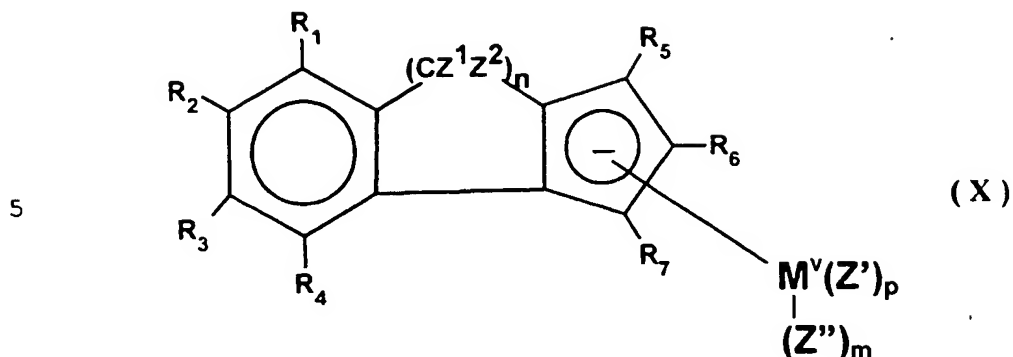
"n" has any of the integer values from 1 to 10 extremes included;

- 20 or a mixture of said compounds having formula (II).

2. The compound according to claim 1, wherein each of the said symbols R_1 , R_2 , R_3 , R_4 , R_5 , R_6 , R_7 , Z^1 and Z^2 represents hydrogen or an alkyl group, linear or branched, having from 1 to 6 carbon atoms, and "n" has values of
- 25 1 to 3.

3. The compound according to any of the previous claims 1 or 2, wherein, in formula (II), Z^1 and Z^2 are both hydrogen.
4. The compound according to claim 3, wherein any two groups selected from those represented by R_1 , R_2 , R_3 and R_4 in formula (II) are linear or branched alkyl having from 1 to 4 carbon atoms.
5. The compound according to any of the previous claims, wherein, in formula (II), the symbols R_1 , R_2 , R_3 , R_4 , Z^1 and Z^2 all represent hydrogen and at least one of the symbols R_5 , R_6 or R_7 , independently represent methyl or ethyl, preferably methyl.
6. The compound according to any of the previous claims, wherein, in formula (II), one of the groups R_5 , R_6 or R_7 , preferably R_5 or R_7 , is a divalent hydrocarbon or silane group having from 2 to 6 carbon atoms, further bound to a second cyclopentadienyl group having from 5 to 20 carbon atoms, preferably selected from cyclopentadienyl, indenyl, fluorenyl and homologous groups thereof.
7. A saline compound or complex comprising at least one polycyclic cyclopentadienyl anion, formally deriving from any of the compounds having formula (II) according to the previous claim 1 by the extraction of an H^+ acid ion from the cyclopentadienyl ring, having the

following general formula (X):



wherein:

each symbol R_1 , R_2 , R_3 , R_4 , R_5 , R_6 , R_7 , Z^1 and Z^2 , and
 10 also the deponent "n", has the same meaning, as the
 corresponding symbol in formula (II) according to any
 of the previous claims 1 to 6;

M^v represents any metal of the periodic table of ele-
 ments having an oxidation state (or valence) "v"
 15 greater than zero;

each Z' is independently a group of an anionic nature
 bound to the metal M^v as anion in an ionic couple or
 with a covalent bond of the " σ " type;

Z'' represents an organic radical having from 5 to 30
 20 carbon atoms, comprising a cyclopentadienyl anion π -
 coordinated to the metal M^v ;

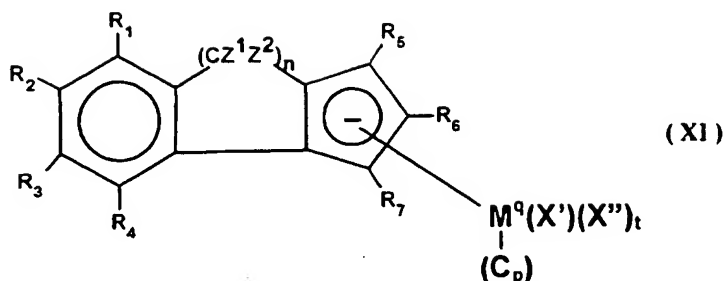
"m" has the value of 1 or 0 depending on whether Z'' is
 present or absent in the compound having formula (X);

"p" has such values that $p = (v - m - 1)$.

25 8. The saline compound according to the previous claim 7,

wherein said metal M^v is selected from metals of groups 1 or 2 of the periodic table.

9. The saline compound according to claim 8, wherein said metal M^v is selected from lithium, sodium or magnesium.
10. The complex according to the previous claim 7, wherein said metal M^v is selected from transition metals of groups 3 to 10 of the periodic table, including metals of the group of lanthanides.
11. The complex according to claims 7 or 10, wherein said metal M^v is selected from metals of groups 4 to 6 of the periodic table, and is preferably Ti, Zr or Hf in oxidation state +4.
12. The complex according to any of the previous claims from 7 to 11, wherein a group selected from said Z' and Z'' in formula (X) is further covalently bound with the polycyclic cyclopentadienyl group deriving from the compounds having formula (II), bound, in the same formula (X), to the metal M^v , by means of a divalent hydrocarbon or silane having from 1 to 15, preferably from 2 to 6, carbon atoms, to form a so-called "bridged" structure.
13. The biscyclopentadienyl complex according to claim 12, having the following general formula (XI):



wherein:

each symbol R_1 , R_2 , R_3 , R_4 , R_5 , R_6 , R_7 , Z^1 and Z^2 , and also the deponent "n", has the same meaning as the corresponding symbol in formula (X) of claim 7;

M^q represents a metal selected from titanium, zirconium and hafnium having an oxidation state (or valence) "q" equal to 3 or 4, preferably 4;

"t" has the value of 1 se the valence "q" of M^q is 4, and the value of 0 if the valence of "q" of M^q is 3;

X' and X'' each independently represent a group of a monovalent anionic nature σ -bound to the metal M^q ;

C_p represents any organic group containing a η^5 -cyclopentadienyl or η^5 -heterocyclopentadienyl ring, of an anionic nature, preferably having at least from 5

to 30 carbon atoms, π -coordinated to the metal M^q .

14. The complex according to claim 13, wherein said symbols X' and X'' each independently represent, a group of an anionic nature σ -bound to the metal M^q selected from hydride, halide, a C_1 - C_{20} alkyl or alkylaryl group, a C_3 - C_{20} alkylsilyl group, a C_5 - C_{20} cycloalkyl

group, a C₆-C₁₀ aryl or arylalkyl group, a C₁-C₁₀ alkoxyl or thioalkoxyl group, a C₂-C₁₀ carboxylate or carbamate group, a C₂-C₁₀ dialkylamide group and a C₁-C₁₀ alkylsilylamide group, or said X' and X'' groups are
5 chemically bound to each other and form a cycle having from 4 to 7 atoms different from hydrogen, also comprising the metal Mⁿ.

15. The complex according to any of the previous claims 13 or 14, wherein said group Cp in the formula (XI) is
10 selected from cyclopentadienyl, indenyl or fluorenyl, and their homologous products, in which one or more carbon atoms of the molecular skeleton, included or not included in the cyclopentadienyl ring, are substituted with a radical selected from the group consisting of chlorine, bromine, a linear or branched alkyl
15 group having from 1 to 10 carbon atoms, optionally halogenated, an alkylsilyl group having from 1 to 10 carbon atoms, a cycloalkyl group having from 5 to 10 carbon atoms, an aryl group having from 6 to 10 carbon
20 atoms, optionally halogenated, an alkoxyl or thioalkoxyl group.

16. The complex according to any of the previous claims 13 or 14, wherein said Cp group in the formula (XI) is
25 selected from anions of the polycyclic cyclopentadienyl compounds having formula (II) according to one of

the claims from 1 to 7, and is preferably equal to the first polycyclic cyclopentadienyl anion of the same formula (XI).

17. The complex according to any of the previous claims 13
5 or 14, wherein said Cp group in the formula (XI) is covalently bound with the polycyclic cyclopentadienyl group deriving from compounds having formula (II), bound, in the same formula (XI), to the metal M^3 , by means of a divalent hydrocarbon or silane group having
10 from 1 to 15, preferably from 2 to 6, carbon atoms, to form a so-called "bridged" structure.

18. A catalyst for the (co)polymerization of ethylene and other α -olefins comprising the following two components in contact with each other, or the product of
15 their reaction:

(i) at least one metallocene complex according to any of the previous claims from 7 to 17, on the condition that the metal M'' in the complex having formula (X) is a metal of groups 4 to 6 of the
20 periodic table;

(ii) a co-catalyst consisting of at least one organo-metallic compound of an element M' different from carbon and selected from elements of groups 2, 12, 13 or 14 of the periodic table as defined
25 above.

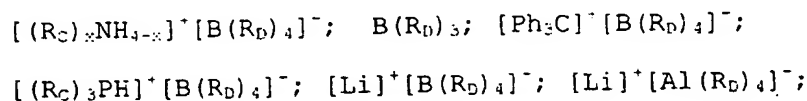
- 19'. The catalyst according to claim 18, wherein said element M' in component (ii) is selected from boron, aluminum, zinc, magnesium, gallium and tin, preferably boron and aluminum.
- 5 20. The catalyst according to one of the claims 18 or 19, wherein said metallocene complex in component (i) is selected from complexes having formula (XI) according to any of the claims from 13 to 17.
21. The catalyst according to any of the previous claims,
10 wherein said component (ii) is a polymeric aluminoxane, preferably methylaluminoxane.
22. The catalyst according to claim 21, wherein the atomic ratio between the transition metal M' in the complex having formula (X) or M³ in the complex having formula
15 (XI), and Al in the aluminoxane ranges from 100 to 5,000.
23. The catalyst according to any of the claims from 18 to 20, wherein said component (ii) consists of at least one compound or a mixture of organometallic compounds
20 of M' capable of reacting with the complex in component (i) by extracting therefrom a σ -bound group to form on the one hand, at least one neutral compound and on the other hand, an ionic compound consisting of a metallocene cation containing the metal M' or M³,
25 and of an organic non-coordinating anion containing

the metal M', whose negative charge is delocalized on a multicentric structure.

24. The catalyst according to claim 23, wherein said M' is B or Al, preferably B.

5 25. The catalyst according to claim 23 or 24, wherein the atomic ratio between the metal M' in component (ii) and the metal M^v or M^z in component (i) ranges from 1 to 6.

26. The catalyst according to any of the previous claims
10 from 23 to 25, wherein said component (ii) consists of an ionic-ionizing compound selected from the group of compounds having one of the following formulae:

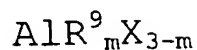


15 wherein the deponent "x" is an integer ranging from 0 to 3;

each R_C group independently represents an alkyl or aryl radical having from 1 to 10 carbon atoms and each R_D group independently represents an aryl radical partially or totally fluorinated, having from 6 to 20
20 carbon atoms.

27. The catalyst according to any of the previous claims from 24 to 26, wherein said component (ii) comprises, in addition to said ionic-ionizing compound, an aluminum trialkyl or an alkylaluminum halide represented
25

by the following formula



wherein: R' is a linear or branched C₁-C₈ alkyl group,
or one of their mixtures,

5 X is a halogen, preferably chlorine or bromine, and

"m" is a decimal number ranging from 1 to 3
extremes included;

with a ratio between M^v or M³ in component (i) and Al
10 in the aluminum alkyl ranging from 1/10 to 1/1,000,
preferably from 1/100 to 1/500.

28. A process for the (co)polymerization of α -olefins,
both in continuous and batchwise, in one or more
steps, at low (0.1-1.0 MPa), medium (1.0-10 MPa) or
15 high (10-150 MPa) pressure, at temperatures ranging
from 20° to 240°C, optionally in the presence of an
inert diluent, characterized in that at least one α -
olefin is put in contact, under the above conditions,
with a catalyst according to one of the previous
20 claims from 18 to 27.

29. The process according to claim 28, wherein said at
least one α -olefin is ethylene.

30. The process according to claim 28 or 29, wherein eth-
ylene is copolymerized with at least a second α -olefin
25 having from 3 to 10 carbon atoms.

31. The process according to claim 30, wherein, in addition to said second α -olefin, ethylene is copolymerized with an aliphatic or alicyclic, non-conjugated diene having from 5 to 20 carbon atoms.
- 5 32. The process according to any of the previous claims from 28 to 31, characterized in that it is carried out in solution or in suspension in a suitable inert liquid medium, consisting of an aliphatic or cycloaliphatic hydrocarbon having from 3 to 15 carbon atoms, or a mixture of these.
- 10 33. The process according to any of the previous claims from 28 to 32, characterized in that the concentration, in the polymerization mixture, either of the metal M^v of the complex having formula (X), or M^d of the complex having formula (XI), forming component (i)
- 15 of said catalyst, ranges from 10^{-5} to 10^{-6} moles/liter.

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INTERNATIONAL SEARCH REPORT

International Application No.

PCT/EP 01/03127

A. CLASSIFICATION OF SUBJECT MATTER IPC 7 C07C13/547 C07C13/54 C07F17/00 B01J31/22 C08F4/42		
According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) IPC 7 C07C C07F B01J C08F		
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched		
Electronic data base consulted during the international search (name of data base and, where practical, search terms used) BEILSTEIN Data, CHEM ABS Data, EPO-Internal		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	CHEMICAL ABSTRACTS, vol. 97, no. 21, 22 November 1982 (1982-11-22) Columbus, Ohio, US; abstract no. 181816s, H. HART ET AL: "1,2-Bridged quadricyclanes" page 770; XP002149350 * compound I * & ISR. J. CHEM, vol. 22, no. 1, 1982, pages 53-55, ---	1
A	EP 0 849 273 A (ENICHEM SPA) 24 June 1998 (1998-06-24) cited in the application claims -----	18,28
<input type="checkbox"/> Further documents are listed in the continuation of box C. <input checked="" type="checkbox"/> Patent family members are listed in annex.		
* Special categories of cited documents : *A* document defining the general state of the art which is not considered to be of particular relevance *E* earlier document but published on or after the international filing date *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) *O* document referring to an oral disclosure, use, exhibition or other means *P* document published prior to the international filing date but later than the priority date claimed *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention *X* document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone *Y* document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art. *G* document member of the same patent family		
Date of the actual completion of the international search 5 June 2001		Date of mailing of the international search report 19/06/2001
Name and mailing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl. Fax: (+31-70) 340-3016		Authorized officer Van Geyt, J

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box I.2

Claims Nos.: 1,3-33 (all in part)

Present claims 1, 7, 18 and 28 relate to an extremely large number of possible compounds. Support within the meaning of Article 6 PCT and/or disclosure within the meaning of Article 5 PCT is to be found, however, for only a very small proportion of the compounds claimed. In the present case, the claims so lack support, and the application so lacks disclosure, that a meaningful search over the whole of the claimed scope is impossible. Consequently, the search has been carried out for those parts of the claims which appear to be supported and disclosed, namely those parts relating to the compounds of claim 2 and their use as catalysts.

The applicant's attention is drawn to the fact that claims, or parts of claims, relating to inventions in respect of which no international search report has been established need not be the subject of an international preliminary examination (Rule 66.1(e) PCT). The applicant is advised that the EPO policy when acting as an International Preliminary Examining Authority is normally not to carry out a preliminary examination on matter which has not been searched. This is the case irrespective of whether or not the claims are amended following receipt of the search report or during any Chapter II procedure.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/LP 01/03127

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP 0849273 A	24-06-1998	IT MI962661 A	19-06-1998
		BR 9705645 A	15-06-1999
		CA 2222917 A	19-06-1998
		CN 1192442 A	09-09-1998
		JP 10231314 A	02-09-1998
		US 5948873 A	07-09-1999
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